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(54) **Heteroatom containing perfluoroalkyl terminated neopentyl glycols and polymers therefrom.**

(57) Heteroatom containing perfluoroalkyl terminated neopentyl glycols of the formulas $\text{HO}[-\text{CH}_2\text{C}(\text{CH}_2\text{X}-\text{E}-\text{R}_1)_2\text{CH}_2\text{O}]_{1-3}-\text{H}$ or $\text{HO}[-\text{CH}_2\text{C}(\text{CH}_2\text{X}-\text{R}_1)_2\text{CH}_2\text{O}]_{1-3}-\text{H}$ are prepared from halogenated neopentyl glycols or oxetanes and thiols of the formula $\text{R}_1-\text{E}-\text{SH}$, amines of the formula $\text{R}_1-\text{E}-\text{NH}-\text{R}$, alcohols of the formula $\text{R}_1-\text{E}-\text{OH}$ or perfluoro-amides, wherein R_1 is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms.

The polymeric reaction products of said fluorinated diols, and optionally other diols with isocyanates to prepare polyurethanes; with amines and isocyanates to prepare polyureas/polyurethanes; with acids or derivatives to prepare polyesters or polycarbonates are disclosed. These include polymeric, random or block urethane, and/or urea, and/or ester compositions containing the residue of at least one R_1 -neopentyl glycol containing two perfluoroalkyl hetero groups.

These polymeric compositions provide improved thermal stability and useful low surface energy oil and water repellent coatings, and soil-release properties for textiles, glass, paper, leather and other materials.

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Description

Heteroatom C ntaining Perfluor alkyl T rminated Neopentyl Glyc ls and Polymers Ther from

This invention relates to hetero group containing perfluoroalkyl terminated neopentyl glycols and their derived polymeric compositions and use to impart oil and water repellency to textiles, glass, paper, leather, and other compositions.

Certain bis-perfluoroalkyl terminated neopentyl glycols devoid of sulfur or nitrogen heteroatoms have been described in U.S. Patent Nos. 3,478,116 and 3,578,701. However, such compounds are difficult and expensive to prepare and therefore are impractical intermediates from which to obtain useful products. Additionally, other bis-perfluoroalkyl glycols containing sulfur are described in U.S. Patent Nos. 3,935,277, 4,001,305, 4,054,592, 4,099,642, and 4,046,944. These compounds are not readily obtainable in high purity and are thermally unstable.

Bis-perfluoroalkyl glycols and polymeric derivatives thereof are useful because they possess a low free surface energy which provides oil and water repellency to a wide variety of substrates. Glycols containing a single R_f -function are known, but do not provide these properties to the same extent. The subject glycols may be prepared in high yield and purity in contrast to prior art materials. Most importantly the instant glycols are thermally stable. Another advantage of the subject glycols is that the pendant perfluoroalkyl chains are connected by flexible hetero groups to the remainder of the molecule thus providing more mobile perfluoroalkyl functions.

One aspect of this invention relates to a method of making such R_f -glycols. Another aspect of this invention relates to derived R_f -containing polyurethane/polyurea/or polyester-containing compositions. One embodiment of this invention relates to hydroxyl terminated prepolymers of the R_f -glycol prepared from polyisocyanates, lactones or isocyanate terminated prepolymers. If desired, R_f -glycols can be converted to isocyanate terminated prepolymers to form reactive compositions with alcohols, polyols, polyamines, R_f -glycols, polyol prepolymers and hydroxyl terminated R_f -containing prepolymers. In another embodiment, the R_f -glycols can be used to replace part or all of the hydroxyl component in a urethane composition.

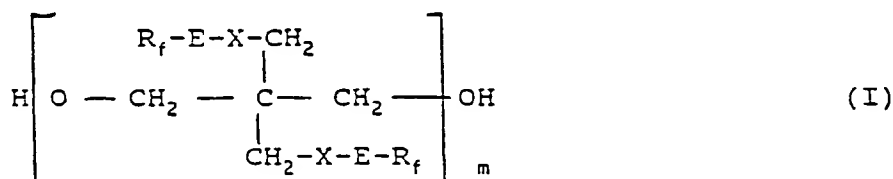
Another embodiment of such polymeric compositions relates to reactive compositions comprising aliphatic polyester segments comprised of the subject R_f -neopentyl glycol reacted with dicarboxylic acids, esters, anhydrides, lactones or acid chlorides, the mole ratios being adjusted to obtain the desired hydroxy or carboxyl end group functionality. In another embodiment suitable functionally terminated polyester polyols are described in Encyclopedia of Polymer Science and Technology, Interscience Pub., 11 (1969), p. 513; these can be linked with the R_f -diol to form the polyesters of this invention. Alternately, the R_f -diol can be coreacted with the polyester precursor reagents, e.g., dicarboxylic acid and aliphatic diol, under esterification conditions. The fluorochemical polyesters can also be derived from isocyanate containing prepolymers, e.g. isocyanate end-capped R_f -glycol and hydroxy terminated polyester segments as described herein.

In another embodiment, the polymers are more appropriately defined as block polymers and contain 15-70% of a fluoro-chemical block connected to 30-85% by weight of a hydrophilic block. These blocks are most preferably connected by urea linkages in the final product condensate and are useful as coatings on textiles, glass, linoleum, leather, wood, tile, metals, plastics, and various materials. They are of particular use on textile materials as soil release agents, showing increased durability as compared to urethane-connected condensation block polymers.

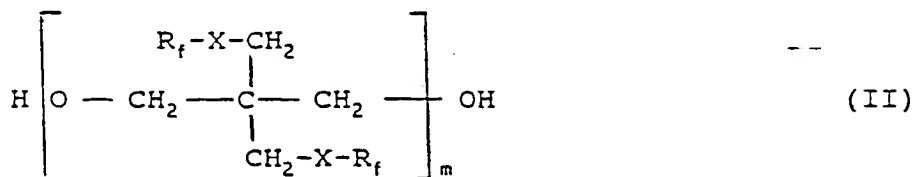
This invention most generally relates to novel heteroatom containing perfluoroalkyl terminated neopentyl glycols and derived polymeric urethane, and/or urea and/or ester linked compositions containing said R_f -glycols.

Another aspect of this invention relates to a substrate containing 0.01 to 10% by weight of a fluorine-containing urethane and/or urea, and/or ester composition, at least part of said fluorine being provided by one or more units derived from the heteroatom containing R_f -neopentyl glycol.

The novel heteroatom containing R_f -neopentyl glycols have the general formulas I or II



or



wherein R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of -NR-, -O-, -S-, -SO₂-, -COO-, -OOC-, -CONR-, -NRCO-, -SO₂NR-, and -NRSO₂-, or said alkylene terminated at the R_f end with -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and for formula I, X is -S-, -O-, -SO₂-, or -NR-, and for formula II, X is -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and where R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms, and m is 1, 2 or 3.

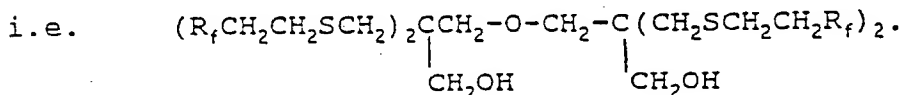
It is understood that the R_f group usually represents a mixture of perfluoroalkyl moieties. When the R_f group is identified as having a certain number of carbon atoms, said R_f group also usually concomitantly contains a small fraction of perfluoroalkyl groups with a lower number of carbon atoms and a small fraction of perfluoroalkyl groups with a higher number of carbon atoms.

Preferably the instant compounds of formula I are those, where R_f is a straight or branched chain perfluoroalkyl of 2 to 12 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, and m is 1 to 2.

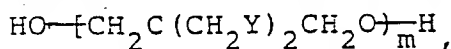
More preferably the instant compounds of formula I are those where R_f is perfluoroalkyl of 2 to 12 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, -CH₂CH₂OCH₂CH₂-, or -SO₂NHCH₂CH₂-, X is -S-, -SO₂- or -O-, and m is 1 or 2.

Most preferred are those compounds where R_f is perfluoroalkyl of 6 to 12 carbon atoms, E is ethylene, m = 1, and X is S, i. e.,
 $(R_fCH_2CH_2SCH_2)_2C(CH_2OH)_2$.

In another group of most preferred compounds R_f is perfluoroalkyl of 6 to 12 carbon atoms, E is ethylene, m = 2, and X is S



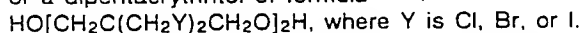
Examples of preferred compounds according to this invention are (C₆F₁₃CH₂CH₂SCH₂)₂C(CH₂OH)₂, (C₈F₁₇CH₂CH₂SCH₂)₂C(CH₂OH)₂, C₇F₁₅CONHCH₂CH₂SCH₂)₂C(CH₂OH)₂, [(CF₃)₂CFO(CF₂)_nCH₂CH₂SCH₂)₂C(CH₂OH)₂, where n is 3 or 4, [(CF₃)₂CFOCF₂CH₂CH₂SCH₂)₂C(CH₂OH)₂, HO[CH₂C(CH₂SCH₂CH₂R_f)₂CH₂O]₂H where R_f is a mixture of C₈F₁₇ and C₁₀F₂₁, (C₈F₁₇CH₂CH₂SO₂CH₂)₂C(CH₂OH)₂, (R_fCH₂CH₂SCH₂)₂C(CH₂OH)₂ where R_f is a mixture of C₄F₉, C₆F₁₃, C₈F₁₇, and C₁₀F₂₁, (R_fSO₂N(C₂H₅)CH₂)₂C(CH₂OH)₂ and (R_fCH₂CH₂OCH₂)₂C(CH₂OH)₂ where R_f is a mixture of C₄F₉, C₆F₁₃, C₈F₁₇ and C₁₀F₂₁. A further object of the invention is a process for the preparation of compounds of the formulas I or II, which comprises reacting compounds of formulas R_f-E-X-H or R_f-X-H, wherein R_f , E and X have the meanings given above, with a halogenated diol of formula HO-



wherein m is 1, 2 or 3 and Y is Cl, Br or I.

The novel R_f -glycols can be obtained directly by the reaction of a perfluoroalkyl thiol of formula R_f-E-SH or a perfluoroalkyl amine of formula R_f-E-NR₂ with a dihalogenated pentaerythritol of formula - (YCH₂)₂C(CH₂OH)₂.

or a dipentaerythritol of formula -



In one preferred embodiment, the neopentyl derivative is dibromopentaerythritol and has the formula (BrCH₂)₂C(CH₂OH)₂. In another preferred embodiment tetrabromodipentaerythritol HO[CH₂C(CH₂Br)₂CH₂O]₂H, is used. These intermediates are commercially available in high purity; dichloro and iodo neopentyl glycols have also been reported.

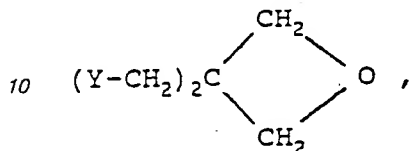
The synthesis of R_f -diols proceeds by the nucleophilic substitution of a perfluoroalkyl substituted thiolate or amine for halide. The reaction may be conducted in an aqueous system using phase transfer catalysis, but work-up of such an aqueous product is difficult due to troublesome emulsions. The improved process of this invention involves the combination of:

- an aprotic solvent, such as N-methylpyrrolidone, N,N-dimethylformamide, dimethyl sulfoxide, or the like, or ketones, such as acetone, methyl alkyl ketones, or dialkyl ketones. Chlorinated solvents and esters generally give poor conversions and are unsuitable;
- moderate reaction temperatures, on the order of 50 to about 140° C; and
- a stoichiometric quantity of an anhydrous alkaline earth carbonate, preferably potassium carbonate, in the ratio of 1 mole of carbonate per mole of halide to be displaced; and
- in the case of the amines, tertiary amine catalysis is useful as exemplified by triethylamine, tributylamine, dimethylaminopyridine, or piperidine.
- in the case of the oxygenated ethers or sulfonamides, Crown ether catalysis is useful as exemplified by 12-Crown-4, 15-Crown-5, and 18-Crown-6.

The reaction temperature, and choice of solvent are mutually dependent. A reaction temperature in the range of 50-140°C is one wherein the formation of undesirable by-products is minimized and wherein the reaction products are stable.

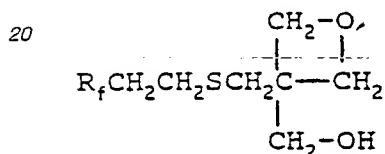
Conditions are adjusted in order to achieve a reasonable rate of reaction at the chosen temperature.

- 5 Alternately, the subject diols are obtained indirectly by the reaction of 3,3-bis-(halomethyl)oxetane of formula



where Y is Cl, Br, or I, with a perfluoroalkanol of formula $\text{R}_f\text{-E-OH}$ or a perfluoroalkylsulfonamide of formula $\text{R}_f\text{-SO}_2\text{NHR}$, followed by hydrolysis of the oxetane linkage to diol.

15 In the synthesis of the R_f -glycols a number of by-products may be present. These are derived in part from trace impurities present in the diol and in part as reaction intermediates. When the starting thiol is $\text{R}_f\text{-CH}_2\text{CH}_2\text{SH}$ and dibromoneopentyl glycol is used these by-products include -



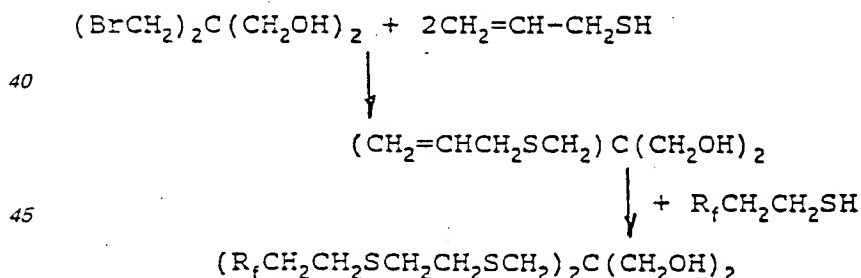
25 $\text{R}_f\text{CH}_2\text{CH}_2\text{SCH}_2\text{C}(\text{CH}_2\text{OH})_3$ and
 $\text{R}_f\text{CH}_2\text{CH}_2\text{SCH}_2\text{C}(\text{CH}_2\text{OH})_2\text{CH}_2\text{Br}$

Such intermediates and the oxetane formation are consistent with the general reaction conditions.

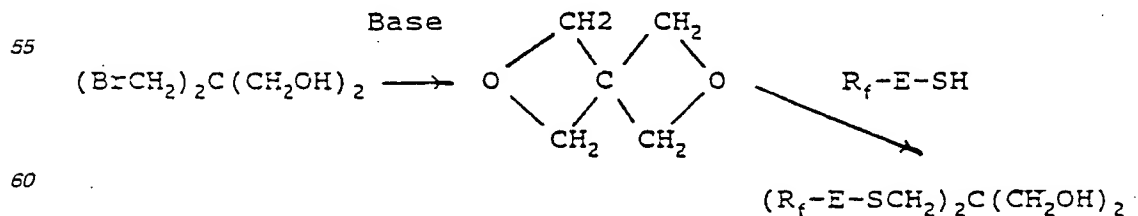
- 30 It should be noted that the ready oxidation of thiols to disulfides requires that the chemistry be conducted in an inert atmosphere.

The subject diols can also be made by first reacting the bromodiols with a functional thiol or amine, e.g. $\text{HSCH}_2\text{CH}=\text{CH}_2$, HSCH_2COOH or $\text{NH}_2\text{CH}_2\text{CH}=\text{CH}_2$. The resultant sulfide or amine can then be reacted with the R_f -containing moiety by a suitable chemistry which does not involve the pendant hydroxyl groups.

- 35 For example,



- 50 An alternate synthesis has also been demonstrated. Dibromoneopentyl glycol can be converted to 2,6-dioxaspiro-[3.3]heptane as reported by Abdun-Nur and Issidorides in J. Org. Chem. 27, 67-69 (1962), and this intermediate converted to R_f -diol.



- 65 This synthetic approach is less favorable since the volatile spiro-oxetane must first be isolated. Attack by the thiolate nucleophile occurs readily on the spiro compound, but the resultant oxetane reacts with difficulty.

The process involves the combination of:

- a. a solvent in which the spiro-oxetane is soluble, preferably n-butanol;
- b. reaction temperatures of 55-150°C. The reaction temperature is kept low initially to avoid volatilization of the oxetane and is increased as the reaction proceeds; and
- c. a basic catalyst, e.g. sodium methoxide or potassium t-butoxide.

The reaction temperature and solvent are mutually dependent. Under appropriate conditions with n-butanol, at 55°C/4.5 hours (1/2 thiol charge) and then 125°C/16.5 hours (remainder of thiol charge), the product formed assayed 17% monoadduct, 78% diadduct.

The subject sulfido-linked diols can be readily oxidized to the corresponding bis-sulfone diols by peracetic acid (H₂O₂/acetic acid) or by other conventional oxidants which selectively oxidize sulfides in the presence of alcohol functions. With peracetic acid, temperatures of 30-100°C are appropriate depending on the amount of excess oxidizing agent to ensure that the intermediate sulfoxides are completely oxidized.

Perfluoroalkyl thiols useful herein are well documented in the prior art. For example, thiols of the formula R_f-E-SH have been described in a number of U.S. Patent Nos. including 3,655,732 and 4,584,143.

Thus, U.S. patent No. 3,655,732 discloses mercaptans of formula

R_f-E-SH

where

E is alkylene of 1 to 16 carbon atoms and R_f is perfluoroalkyl, and teaches that halides of formula R_f-E-Halide are well-known; reaction of R_fI with ethylene under free-radical conditions gives R_f(CH₂CH₂)_nI while reaction of R_fCH₂I with ethylene gives R_fCH₂(CH₂CH₂)_nI as is further taught in U.S. Patent Nos. 3,088,849; 3,145,222; 2,965,659 and 2,972,638.

U.S. Patent No. 3,655,732 further discloses compounds of formula R_f-R'-Y-R''-SH

where

R' and R'' are alkylene of 1 to 16 carbon atoms, with the sum of the carbon atoms of R' and R'' being no greater than 25; R_f is perfluoroalkyl of 4 through 14 carbon atoms and Y is -S- or -NR'''- where R''' is hydrogen or alkyl of 1 through 4 carbon atoms.

U.S. Pat. No. 3,544,663 teaches that the mercaptan

R_fCH₂CH₂SH

where

R_f is perfluoroalkyl of 5 to 13 carbon atoms, can be prepared by reacting the perfluoroalkyl alkylene iodide with thiourea or by adding H₂S to a perfluoroalkyl substituted ethylene (R_f-CH=CH₂), which in turn can be prepared by dehydrohalogenation of the halide R_f-CH₂CH₂- halide.

The reaction of the iodide R_f-E-I with thiourea followed by hydrolysis to obtain the mercaptan R_f-E-SH is the preferred synthetic route. The reaction is applicable to both linear and branched chain iodides.

Particularly preferred herein are the thiols of formula

R_fCH₂CH₂SH

where

R_f is perfluoroalkyl of 6 to 12 carbon atoms. These R_f-thiols can be prepared from R_fCH₂CH₂I and thiourea in very high yield.

Perfluoroalkylamines useful herein are well documented in the prior art. For example, C₆F₁₃CH₂CH₂NH₂ has been described in Japan Kokai 77/118,406. R_fCH₂NH₂ wherein R_f is CF₃ through CF₃(CF₂)₁₁ are described in British patent No. 717,232 (1954).

Further R_fSO₂NR(CH₂)_nNR(CH₂)₃NH₂ and R_fCH₂CH₂SO₂NH(CH₂)_nNR₂ are described in G.B. 1,106,641 and U.S. 3,838,165 respectively; R_fCONH(CH₂)_nNH₂ in Jap. Kokai 52/14767.

Perfluoroalkanols useful herein are well documented in the prior art and many are commercially available. They have the general formula R_f-E-OH and include:

C₈F₁₇SO₂N(C₂H₅)C₂H₄OH

C₈F₁₇C₂H₄OH

C₇F₁₅CH₂OH

C₇F₁₅CON(C₂H₅)C₂H₄OH

C₈F₁₇C₂H₄SO₂C₂H₄OH

(CF₃)₂CF(CF₂)₈C₂H₄OH

(CF₃)₂CFOC₂F₄C₂H₄OH

C₈F₁₇C₂H₄SO₂N(CH₃)C₄H₈OH

C₈F₁₇CH₂OH

CF₃CH₂OH

C₈F₁₇CH₂CH₂SO₂NHCH₂CH₂OH

Perfluoroalkylsulfonamides useful herein are well documented in the prior art such as in U.S. 2,915,554 and include compounds of the general structure R_f-SO₂NHR, such as--

C₈F₁₇SO₂N(C₂H₅)OH

C₈F₁₇SO₂N(CH₃)H

C₈F₁₇SO₂N(i-C₃H₇)H

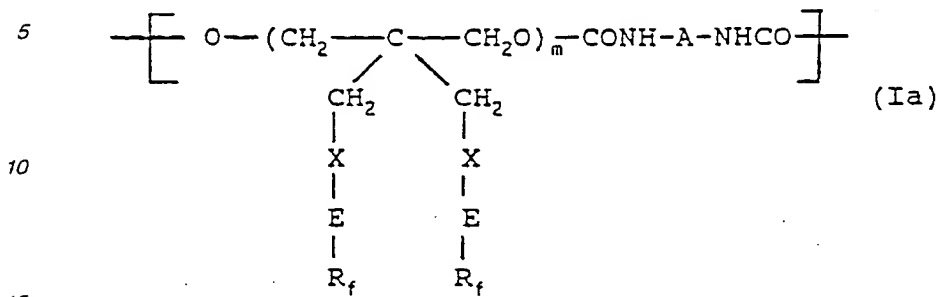
C₁₀F₂₁SO₂N(C₂H₅)H

C₁₀F₂₁SO₂NH₂

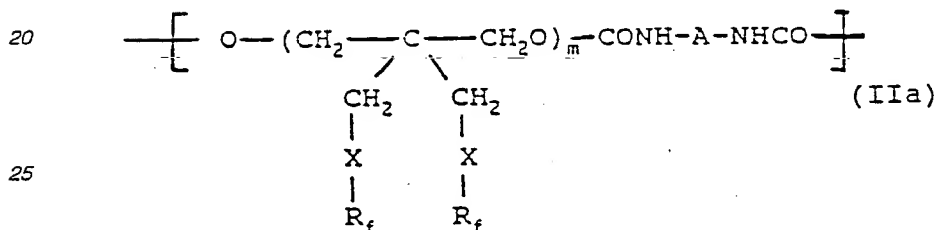
The diols, can be used directly or indirectly to make a variety of polymers such as polyesters, polyureas,

polycarbonates, polyurethanes and the like. Polyurethanes are particularly preferred.

A further object of the invention are linear polyurethanes having recurring structural units of the formula Ia or IIa:



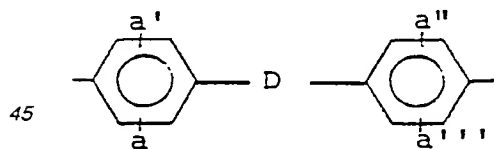
or



wherein R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of $-\text{NR}-$, $-\text{O}-$, $-\text{S}-$, $-\text{SO}_2-$, $-\text{COO}-$, $-\text{OOC}-$, $-\text{CONR}-$, $-\text{NRCO}-$, $-\text{SO}_2\text{NR}-$, and $-\text{NRSO}_2-$, or terminated at the R_f end with $-\text{CONR}-$ or $-\text{SO}_2\text{NR}-$, where R_f is attached to the carbon or sulfur atom, and for formula Ia, X is $-\text{S}-$, $-\text{O}-$, $-\text{SO}_2-$, or $-\text{NR}-$, and for formula IIa, X is $-\text{CONR}-$ or $-\text{SO}_2\text{NR}-$, where R_f is attached to the carbon or sulfur atom, and where R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms, and m is 1, 2 or 3, and A is a divalent aliphatic, cycloaliphatic or aromatic group.

Preferably A is alkylene of 2 to 16 carbon atoms, cycloalkylene of 6 to 15 carbon atoms, phenylene, naphthylene or said phenylene or said naphthylene substituted by one or two alkyl of 1 to 4 carbon atoms, by alkoxy of 1 to 4 carbon atoms, by chloro, by bromo or by nitro, or A is a group of the formula



where D is a direct bond, oxygen, methylene or ethylene, and a, a', a'' and a''' each independently are hydrogen, alkyl of 1 to 4 carbon atoms, alkoxy of 1 to 4 carbon atoms, chloro or bromo.

More preferably A is 3,3,4-trimethyl-1,16-hexanediyl or 1,3,3-trimethyl- α ,5-hexahydrobenzylene.

In a preferred embodiment, R_f is perfluoroalkyl of 2 to 12 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is alkylene of 2 to 6 carbon atoms, $-\text{CONHCH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{N}(\text{CH}_3)\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{SO}_2\text{NHCH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{OCH}_2\text{CH}_2-$, or $-\text{SO}_2\text{NHCH}_2\text{CH}_2-$, X is $-\text{S}-$, $-\text{SO}_2-$ or $-\text{O}-$, and m is 1 or 2.

Preferred urethane compositions include those where R_f and E have the configurations previously described as being preferred and X is S, SO_2 or O.

The R_f -glycols can be used to make a wide variety of urethane intermediate and end products including hydroxyl and isocyanate-terminated prepolymers, low molecular weight urethane compositions useful to render plastics soil repellent, and high molecular weight compositions useful as elastomers, foams, paints and varnishes, and textile treating compositions. It is also possible to modify these R_f -containing urethane compositions so that they are water soluble or self-emulsifiable, a property that is particularly useful in connection with the textile treating compositions. Block polymers compositions of the R_f containing diols and hydrophilic blocks have special utility in the textile industry as soil-release agents.

These compositions have extremely low free surface energies and therefore, possess oil and water repellent properties, as well as mold release and other properties associated with low free surface energy. It should be

noted that the compositions of this invention are characterized by the presence of two perfluoroalkylhetero groups in close proximity, a characteristic which provides improved oil and water repellent properties over the fluorinated compositions of the prior art. Further the two perfluoroalkylthio groups are connected via a neopentyl moiety which does not permit the thermal elimination of mercaptan by beta-elimination. Hence, these R_f -diols and derivatives have enhanced thermal stability. The beta-elimination reaction of prior art sulfur containing R_f -diols is disclosed in U.S. Patent No. 4,097,642.

Using the R_f -compounds and compositions described herein, it is possible to manufacture molds that display the excellent release properties characteristic of silicone polymers. It is also possible to prepare polymeric compositions with enhanced thermal stability.

Polyurethane elastomers generally have remarkable resistance to most solvents including gasoline, aliphatic hydrocarbons and, to some degree, aromatic hydrocarbons. They also exhibit excellent abrasion resistance. By inclusion of the R_f -glycol in an elastomer formulation, it is possible to increase the solvent resistance of urethane elastomers. The elastomers generally involve the reaction product of a diisocyanate, a linear long chain diol and a low molecular weight chain extender such as a glycol, diamine or polyol. Today, elastomers are generally prepared by a prepolymer technique whereby a diisocyanate is reacted with a hydroxyl-terminated polyester or polyether to form an isocyanato-terminated prepolymer. This prepolymer is then further reacted (chain extended) with a glycol, diamine or polyfunctional polyol (e.g. trimethylolpropane). Following the chain extension step, the liquid material solidifies and is removed from a mold and cured at elevated temperatures.

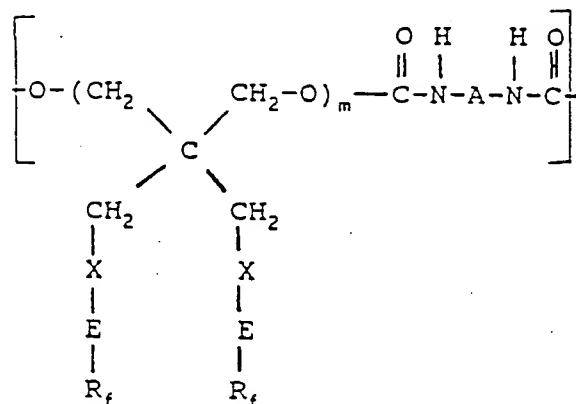
Urethane foams are usually prepared from diisocyanates and hydroxyl-terminated polyethers or polyesters. Linear or slightly branched polymers are used to provide flexible foams while more highly branched polymers produce rigid foams. Foaming is often accomplished by including water in the system, the reaction between isocyanate and water providing carbon dioxide for foaming. For rigid foams a low-boiling liquid such as trichlorofluoromethane has been used as a blowing agent.

Appropriate selection of catalysts, stabilizers, surfactants and other additives controls the foam formation, cell size and type, density, cure and the like. By incorporating the R_f -glycol into urethane foams, especially molded foams, it is possible to achieve improved mold release properties in rigid, semi-rigid and flexible foams. It is also possible to improve the water and solvent resistance of foams used as insulation.

Incorporation of the R_f -glycols into polyurethane coatings such as paints and varnishes improves the water and solvent resistance thereof. Widely used systems include the two-component coatings wherein a non-volatile isocyanate derived from the reaction of tolylene diisocyanate with a polyol, such as trimethylolpropane, is reacted with a polyfunctional polyester. Another system in use involves the one-component polyurethane coatings which are based on stable isocyanate-terminated prepolymers obtained from a diisocyanate such as tolylene diisocyanate and a polyfunctional polyether. Such coatings dry by the reaction of the free isocyanate groups with water or atmospheric moisture. The reaction proceeds through the unstable carbamic acid, with CO_2 being eliminated, to give primary amine groups which further react with isocyanate groups to form ureas.

Treatment of a textile with a fluorine-containing composition, notably a fluorine-containing polymer, provides oil and water-repellent characteristics thereto. Such compositions containing the residue of the R_f -glycol display improved oil and water repellence on textile substrates.

In higher molecular weight urethane compositions, linear polymer, obtained by reacting an R_f -glycol with an organic diisocyanate, having recurring structural units of formula



where

R_f , E and X are as previously defined, where m is 1 or 2, and a similar structure can be drawn for a type (II) diol, and A is a divalent organic radical, preferably alkylene of 2 to 16 carbon atoms, cycloalkylene of 6 to 15 carbon atoms, unsubstituted or substituted phenylene or naphthylene or unsubstituted or substituted biphenylene or bisphenylene. The polymers are useful as plastics, fibers, coatings and the like.

The treatment or modification of fabrics to improve their properties is routine practice in the textile industry.

However, resin-treated durable press garments are difficult to clean because they are prone to soil retention. Investigation of this phenomenon showed that resin treated cotton as well as the synthetic fibers are oleophilic and accordingly, dirt, particularly oily stains, clings tenaciously to the fabric substrate and is extremely difficult to remove under normal home washing conditions.

In an effort to overcome the soil removal resistance of resin-treated fabrics, artisans have commonly treated such fabrics with a hydrophilic colloid such as carboxymethyl cellulose and synthetic polymers such as polyacrylic acid and copolymers of acrylic acid with lower alkyl acrylates and methacrylates. These materials, which are referred to as soil release agents, apparently coat the textile fibers with a hydrophilic film which allows the fibers to be wet effectively by detergent solutions so that the soils are readily removed by laundering.

The combination of oleophobic fluorinated groups and hydrophilic poly(ethyleneoxide) containing groups in one polymer to achieve the release of oil stains from textiles has been described in U.S. patent Nos. 3,728,151 and 3,758,447. The advantage of fluorinated soil release agents over non-fluorinated ones stems from their oleophobic nature, which (a) prevents the wicking of oil stains into the fabrics and (b) facilitates the lifting off of the staining material from the fabric when it is washed.

It has also been reported that certain segmented (block) polyurethane/polyureas are useful soil-release agents - see U.S. Patent No. 4,046,944.

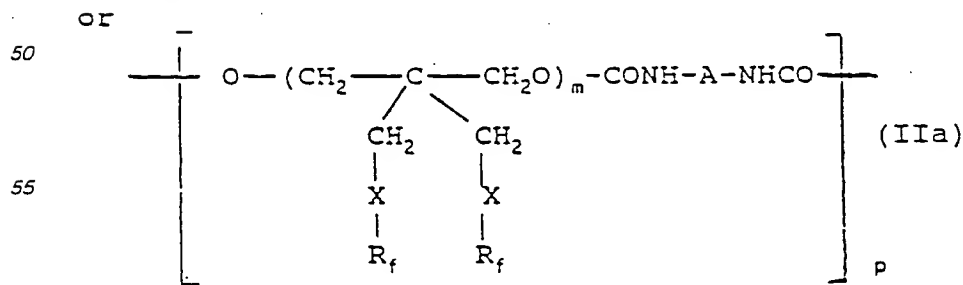
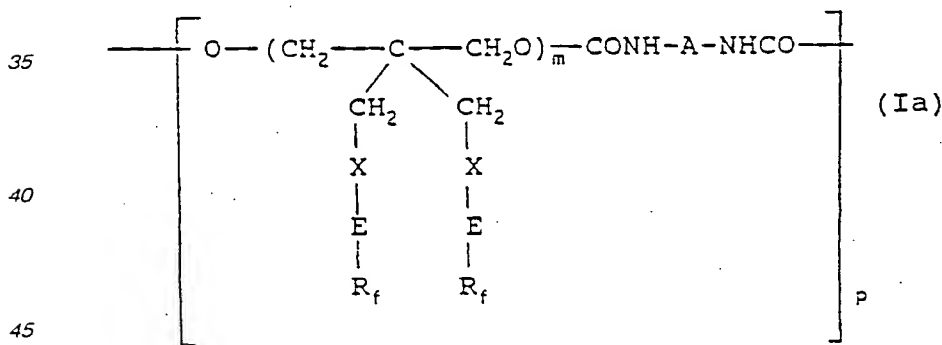
The oleophobic blocks are comprised of fluorinated aliphatic groups held together in segments directly or through linkages made up of various combinations of functional groups and/or hydrocarbon chains. The preferred hydrophilic segments are based on polyoxyalkylene glycols or end-capped derivatives thereof. The molecular weight of the glycols may range from about 150 to 10,000 or more and may be repeated from 1 to 500 or more times.

A preferred embodiment of this invention is that a great increase in performance, especially in durability of fluorochemical textile finishes, can be achieved by use of polymer condensates, wherein the oleophobic and hydrophilic blocks are preferably connected by urea linkages rather than urethane linkages. A block polymer comprising oleophobic fluorinated blocks and hydrophilic poly(ethyleneoxide) blocks connected by urea linkages and having increased durability when applied to materials such as textiles is thus provided.

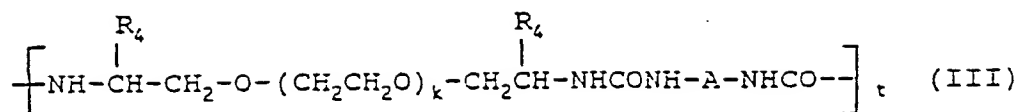
A further embodiment of the invention is a linear block copolymer having soil-release properties which contains, based on the total block copolymer,

(a) 15-70% by weight of recurring structural units of blocks Ia, IIa or a mixture thereof,

(b) 85 to 30% by weight of recurring structural units of block III, wherein the blocks are connected to one another via urea linkages, blocks Ia or IIa having recurring structural units of the formula



block III having recurring structural units of the formula:



wherein R_1 is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of $-\text{NR}-$, $-\text{O}-$, $-\text{S}-$, $-\text{SO}_2-$, $-\text{COO}-$, $-\text{OOC}-$, $-\text{CONR}-$, $-\text{NRCO}-$, $-\text{SO}_2\text{NR}-$, and $-\text{NRSO}_2-$, or terminated at the R_1 end with $-\text{CONR}-$ or $-\text{SO}_2\text{NR}-$, where R_1 is attached to the carbon or sulfur atom, and for formula Ia, X is $-\text{S}-$, $-\text{O}-$, $-\text{SO}_2-$, or $-\text{NR}-$, and for formula IIa, X is $-\text{CONR}-$ or $-\text{SO}_2\text{NR}-$, where R_1 is attached to the carbon or sulfur atom, and where R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms, and m is 1, 2 or 3.

A is a divalent aliphatic, cycloaliphatic or aromatic group,

R_4 is hydrogen or methyl,

k is 8 to 100, and

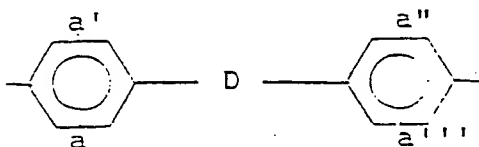
p and t are integers and are independently 1 to 5.

It will be noted that the invention condensation polymers, being made up of Blocks Ia and/or IIa and III are of the pattern:

(oleophobic R_1 -block)-NH-C(=O)-NH-[hydrophilic poly(ethylene oxide) block]

Preferably R_1 is perfluoroalkyl of 2 to 12 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is alkylene of 2 to 6 carbon atoms, $-\text{CONHCH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{N}(\text{CH}_3)\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{SO}_2\text{NHCH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{OCH}_2\text{CH}_2-$, or $-\text{SO}_2\text{NHCH}_2\text{CH}_2-$, X is $-\text{S}-$, SO_2- or $-\text{O}-$, and m is 1 or 2.

Preferably A is alkylene of 2 to 16 carbon atoms, cycloalkylene of 6 to 15 carbon atoms, phenylene, naphthylene or said phenylene or said naphthylene substituted by one or two alkyl of 1 to 4 carbon atoms, by alkoxy of 1 to 4 carbon atoms, by chloro, by bromo or by nitro, or A is a group of the formula



where D is a direct bond, oxygen, methylene or ethylene, and a, a', a'' and a''' each independently is hydrogen, alkyl of 1 to 4 carbon atoms, alkoxy of 1 to 4 carbon atoms, chloro or bromo.

More preferably A is 3,3,4-trimethyl-1,6-hexanediyl of 1,3,3-trimethyl- α ,5-hexahydrobenzyiene.

The molecular weight of Blocks Ia, IIa and III will vary, depending on the substituents and the number of repeating units in each. However, the advantages combination of Blocks Ia and/or IIa and III can be achieved by the use of monomer or prepolymer reactants so as to give 15-70 % Block I to 30 % to 85 % Block II in the final polymer:

The segmented perfluoroalkyl/hydrophilic polymers are useful on substrates as coatings, which will (1) prevent, or at least reduce, soiling and (2) release soil when washed with water. They are therefore useful as ingredients in floor polishes, furniture waxers, window washing fluids, and so on; their most important application is as a soil-release finish on textiles, especially polyester/cotton textiles. Generally, they are useful as coatings on glass, ceramics, masonry, wood, plastics, textiles, leather and metals, or as additive ingredients in such coatings.

Most urethane compositions that are used commercially to any great extent are copolymers that contain only a relatively small number of urethane linkages. These copolymers are prepared from a variety of segments, typically based on polyethers and polyesters and can have a molecular weight of from 200 to 10,000, generally from about 200 to about 4,000. By the inclusion of an appropriate amount of R_1 -glycol in the starting materials, it is possible to prepare prepolymers that, when incorporated as part of a urethane composition favorably affect the properties thereof. It is similarly possible to incorporate a desired amount of R_1 -glycol into the reaction mixture of a conventional prepolymer and an isocyanate so as to obtain conventional urethane compositions containing the divalent residue of the R_1 -glycol. In the same way, there can be added an R_1 -containing prepolymer together with or instead of the R_1 -glycol.

The R_1 -containing prepolymers can be hydroxy-terminated or isocyanate-terminated and, as indicated, can have a molecular weight as high as 10,000 although a molecular weight of 200 to about 4,000 is more usual.

Hydroxy-terminated prepolymers can be prepared by reacting an excess of a polyhydroxy component with a polyfunctional hydroxy-reactive component such as a polyisocyanate; an isocyanate-terminated prepolymer; a polybasic carboxylic acid, anhydride or acrylyl halide; phosgene; or a bischloroformate.

The polyhydroxy component can be a polyol, an R_1 -glycol, a polyether, a polyester, an R_1 -containing polyether, an R_1 -containing polyester or mixture thereof.

The polyols are well-known in the urethane art and include:

Ethylene glycol, 1,3-propanediol, 1,4-butanediol, 1,5-pentanediol, 1,6-hexanediol, 1,9-nonanediol, 1,10-decanediol, di-, tri-, tetra- and pentaethylene glycol, bis(4-hydroxybutyl) ether, bis(2-hydroxyethyl) thioether, bis(4-hydroxybutyl) thioether, 1,4-bis(3-hydroxypropyl)benzene, glycerol, trimethylolpropane, 1,2,6-hexanetriol, sorbitol, mannitol, pentaerythritol, 2-ethyl-1,3-butylene glycol, octamethylene glycol, 2-ethyl-1,3-hexanediol, dodecamethylene glycol, tetradecamethylene glycol, hexadecamethylene glycol, octadecamethylene glycol.

The polyol can also contain cycloaliphatic groups, e.g. 1,4-cyclohexane-diol, 1,4-bis(hydroxymethyl)cyclohexane, 4,4'-dihydroxyl-1,1'-dicyclohexyl and the like. If desired, mixtures of polyols can be used.

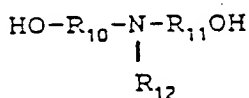
Polyols in addition to those described above, that are considered especially useful, are those containing tertiary nitrogen atoms which can be quaternized with acids, thereby converting a water-insoluble urethane composition into one that is water soluble or emulsifiable. Generally, an isocyanate-terminated prepolymer having a molecular weight of 200 to 10,000, preferably 400 to 4,000, is reacted with a difunctional tertiary amine to provide a segmented polymer containing tertiary nitrogen atoms. The nitrogen atoms can be quaternized, for example, by alkylation with methyl chloride or dimethyl sulfate to yield a composition that in polar media yields a dispersion in water. The polyammonium polyurethane compositions are obtained even more readily by neutralization of the basic polyurethane composition in a polar organic solvent such as acetone, methyl ethyl ketone, tetrahydrofuran, with a strong (HCl) or preferably weak ($pK > 4$) acid such as the C_2 - C_9 alkanolic acids. Acetic acid is especially preferred because the acetic acid evaporates with the water on drying to leave the water-insoluble hydrophobic starting polyurethane composition.

The neutralized polyurethane composition in a polar solvent spontaneously forms a dispersion when water is added. The solvent can thereafter be distilled off to give a solvent-free latex whose film-forming qualities are comparable to those of the organic solution.

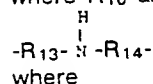
In a convenient mode of preparing the water-dispersible basic polyurethane compositions, a polyester or polyether diol is reacted in a non-reactive polar solvent, such as acetone, methyl ethyl ketone, tetrahydrofuran and the like, with an excess of a diisocyanate such as tolylene diisocyanate or, preferably an aliphatic diisocyanate which tends to give non-yellowing urethanes such as dimer acid derived diisocyanate (DDI, commercially available from Quaker Oats Company) or another diisocyanate which is described herein as providing non-yellowing urethanes, and the prepolymer partially chain extended with an alkyl diethanolamine to yield a urethane composition containing tertiary amino groups. The urethane composition can then be acidified with a solution of aqueous weak acid ($pK > 4$) such as acetic acid; the concentration of acid is not critical. An emulsion immediately forms when this composition is added to water.

The polyurethane compositions can contain from as little as 5 to 800 milliequivalents of ammonium groups per 100 grams of polyurethane composition, preferably from about 50 to about 500 milliequivalents of ammonium groups per 100 grams.

Some useful polyols containing tertiary nitrogen atoms can be represented by the formula:



where R_{10} and R_{11} are alkyl of 2 to 4 carbon atoms or a group of formula



R_{13} and R_{14} are alkyl of 2 to 4 carbon atoms

R_{12} is alkyl of 1 to 18 carbon atoms, cyclohexyl, tolyl, xylyl, naphthyl, or pyridyl.

Useful polyols that contain tertiary nitrogen atoms include the alkoxyated aliphatic, cycloaliphatic aromatic and heterocyclic primary amines:

N-methyl-diethanolamine, N-butyl-diethanolamine, N-oleyl-diethanolamine, N-cyclohexyl-diethanolamine, N-methyl-diisopropanolamine, N-cyclohexyl-diisopropanolamine, N,N-dihydroxyethylaniline, N,N-dihydroxyethyl-m-toluidine, N,N-dihydroxyethyl-p-toluidine, N,N-dihydroxypropyl-naphthylamine, N,N-tetrahydroxyethyl-aminopyridine, polyethoxylated butyldiethanolamine, polypropoxylated methyldiethanolamine (molecular wt. 1000), and polypropoxylated methyldiethanolamine (molecular wt. 2000); also useful are polyesters with tertiary amino groups, tri-2-hydroxypropyl-(1)-amine, N,N-di-n-(2:3-dihydroxypropyl)-amine, N,N'-bis-hydroxypropylethylenediamine, N,N'-dimethyl-N,N'-bis-(hydroxyethyl)-ethylenediamine, 11-stearyldiethanolamine.

The R_i -glycols can be incorporated in the water-dispersible urethane compositions in an amount sufficient to provide the desired improvement in the surface properties of the polyurethane composition.

Useful polyethers are well-known and widely employed in urethane technology.

The polyethers are generally prepared commercially from lower alkylene oxides e.g. ethylene, propylene and butylene oxide and di- or polyfunctional alcohols. They have a molecular weight of from 400 to 5000. A list of commercially available polyethers, trade names, molecular weight range and suppliers can be found in Volume 11, Polyurethane, page 511, Encyclopedia of Polymer Science and Technology, John Wiley and Sons, Inc., 1969.

A further embodiment of the invention is a linear polyester having recurring structural units of the formula Ib or IIb

--[OCH₂C(CH₂X-E-R₁)₂CH₂O]_m-CO-L-CO]- (Ib)

or

--[OCH₂C(CH₂X-R₁)₂CH₂O]_m-CO-L-CO]- (IIb)

wherein R₁ is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of -NR-, -O-, -S-, -SO₂-, -COO-, -OOC-, -CONR-, -NRCO-, -SO₂NR-, and -NRSO₂-, or terminated at the R₁ end with -CONR- or -SO₂NR-, where R₁ is attached to the carbon or sulfur atom, and for formula Ib, X is -S-, -O-, -SO₂-, or -NR-, and for formula IIb, X is -CONR- or -SO₂NR-, where R₁ is attached to the carbon or sulfur atom, and where R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms, and m is 1, 2 or 3.

L is a direct bond, a straight or branched chain alkylene of 1 to 34 carbon atoms, cycloalkylene of 6 to 12 carbon atoms, or arylene of 6 to 16 carbon atoms.

Preferred polyesters are those where R₁ is perfluoroalkyl of 2 to 12 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, -CH₂CH₂OCH₂CH₂CH₂-, or -SO₂NHCH₂CH₂-, X is -S-, SO₂- or -O-, and m is 1 or 2.

Also preferred are polyesters wherein L is tetramethylene or o-phenylene.

Hydroxy-terminated polyesters can be prepared from a polybasic acid, anhydride or aryl halide and a polyol, as described above and/or an R₁-glycol.

Useful dicarboxylic acids are those derived from a saturated aliphatic or cycloaliphatic dicarboxylic acid of 2 to 36 carbon atoms or an aromatic dicarboxylic acid of 8 to 18 carbon atoms, e.g. compounds of formula L(COOH)₂ where L is preferably alkylene of 0-16 carbon atoms or arylene of 6 to 16 carbon atoms. Such acids include oxalic, malonic, succinic, glutaric, adipic, pimelic, suberic, azelaic, sebacic, brassylic, octadecanedioic, dimer acid, 1,4-cyclohexanedicarboxylic, 4,4'-dicyclohexyl-1,1'-dicarboxylic, phthalic, isophthalic, terephthalic, methylphthalic, chlorophthalic, diphenyl-2,2'-dicarboxylic, diphenyl-4,4'-dicarboxylic, 1,4-naphthalene dicarboxylic, diphenylmethane-2,2'-dicarboxylic, diphenylmethane-3,3'-dicarboxylic, diphenylmethane-4,4'-dicarboxylic acid and the like.

Adipic acid and phthalic anhydride are the most common acid and anhydride. Of the polyols, the most commonly used include ethylene glycol, propylene glycol, 1,2-, 1,3- and 1,4-butylene glycol, 1,6-hexylene glycol, trimethylolpropane, glycerol, 1,2,6-hexanetriol and diethylene glycol.

Useful hydroxyl-terminated polyesters can also be derived from natural castor oil and glycerol, from caprolactones and ethylene glycol. Such hydroxy-terminated polyesters have hydroxyl numbers ranging from 40 to 500 and very low acid numbers ranging from 0 to 2.

Hydroxyl-terminated polycarbonates can be obtained by reacting an excess of a polyol with phosgene. Hydroxyl terminated polyether carbonates are also available and suitable for this invention.

Hydroxy-terminated polybutadienes, or butadiene styrenes and butadiene-acrylonitriles are useful herein, as are hydroxyl containing graft polymers of the polyether polyacrylonitrile type.

Any convenient isocyanate can be used to react with the R₁-glycol or R₁-containing hydroxy-terminated prepolymer. Many useful isocyanates are well-known in the art. Thus, one can use aliphatic or aromatic isocyanates, diisocyanates, triisocyanates and polyisocyanates.

Useful aromatic diisocyanates can be represented by the formula

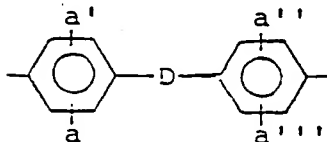
A(NCO)₂

where

A is phenylene that is unsubstituted or substituted by one or two alkyls of 1 to 4 carbon atoms, alkoxy of 1 to 4 carbon atoms, chloro, bromo and nitro, naphthylene that is unsubstituted or substituted by one or two of alkyl of 1 to 4 carbon atoms, chloro, bromo and nitro

or where

A is a group of formula



where

D is a direct bond, oxygen, methylene or ethylene and a, a', a'' and a''' each independently are hydrogen, alkyl of 1 to 4 carbon atoms, alkoxy of 1 to 4 carbon atoms, chloro or bromo.

Aromatic triisocyanates can be represented by the formula G(NCO)₃

where

G is the benzene or toluene group.

Aromatic di- and triisocyanates as described above include

Toluene diisocyanate (TDI) (all isomers), 4,4'-diphenylmethane diisocyanate (MDI), toluene diisocyanate, dianisidine diisocyanate, m-xylene diisocyanate, p-phenylene diisocyanate, m-phenylene diisocyanate,

1-chloro-2,4-phenylene diisocyanate, 3,3'-dimethyl-4,4'-bisphenylene diisocyanate, 4,4'-bis(2-methylisocyanatophenyl)methane, 4,4'-bisphenylene diisocyanate, 4,4'-bis(2-methoxyisocyanatophenyl)methane, 1-nitrophenyl-3,5-diisocyanate, 4,4'-diisocyanatodiphenyl ether, 3,3'-dichloro-4,4'-diisocyanatodiphenyl ether, 3,3'-dichloro-4,4'-diisocyanatodiphenyl-methane, 4,4'-diisocyanatodibenzyl, 3,3'-dimethoxy-4,4'-diisocyanatodiphenyl, 2,2'-dimethyl-4,4'-diisocyanatodiphenyl, 2,2'-dichloro-5,5'-dimethoxy-4,4'-diisocyanatodiphenyl, 3,3'-dichloro-4,4'-diisocyanatodiphenyl benzene-1,2,4-triisocyanate, benzene-1,3,5-triisocyanate, benzene-1,2,3-triisocyanate, toluene 2,4,6-triisocyanate, toluene 2,3,4-triisocyanate, 1,2-naphthalene diisocyanate, 4-chloro-1,2-naphthalene diisocyanate, 4-methyl-1,2-naphthalene diisocyanate, 1,5-naphthalene diisocyanate, 1,6-naphthalene diisocyanate, 1,7-naphthalene diisocyanate, 1,8-naphthalene diisocyanate, 4-chloro-1,8-naphthalene diisocyanate, 2,3-naphthalene diisocyanate, 2,7-naphthalene diisocyanate, 1,8-dinitro-2,7-naphthalene diisocyanate, 1-methyl-2,4-naphthalene diisocyanate, 1-methyl-5,7-naphthalene diisocyanate, 6-methyl-1,3-naphthalene diisocyanate, 7-methyl-1,3-naphthalene diisocyanate, polymethylene polyphenyl isocyanate and co-products of hexamethylene diisocyanate and tolylene diisocyanate.

Useful aliphatic diisocyanates include those of general formula

$A(NCO)_2$

where

A is straight or branched chain alkylene of 2 to 16 carbon atoms, optionally containing halides or cycloaliphatic functions.

Useful aliphatic or cycloaliphatic polyisocyanates include -1,2-ethane diisocyanate, 1,3-propane diisocyanate, 1,4-butane diisocyanate, 2-chloropropane-1,3-diisocyanate, pentamethylene diisocyanate, propylene-1,2-diisocyanate, 1,6-hexane diisocyanate, 1,8-octane diisocyanate, 1,10-decane diisocyanate, 1,12-dodecane diisocyanate, 1,16-hexadecane diisocyanate and other aliphatic diisocyanates such as 1,3-cyclohexane diisocyanate, 1,4-cyclohexane diisocyanate, cyclohexane triisocyanate, 4,4'-methylene bis(cyclohexyl isocyanate).

Additionally, the following diisocyanates are particularly preferred because urethane compositions made therefrom tend to be non-yellowing:

1,6-hexamethylene diisocyanate (HDI), 2,2,4- and 2,4,4-trimethylhexamethylene diisocyanate (TMDI), dimer acid derived diisocyanate (DDI) obtained from dimerized fatty acids, such as linoleic acid 4,4'-dicyclohexylmethane diisocyanate (hydrogenated MDI), isophorone diisocyanate, 3-isocyanatomethyl-3,5,5-trimethylcyclohexyl diisocyanate, lysine methyl ester diisocyanate (LDIM), bis(2-isocyanatoethyl) fumarate (FDI), bis(2-isocyanatoethyl) carbonate, m-tetramethylxylene diisocyanate (TMXDI), additional polyisocyanates include polymethylene polyphenylisocyanate (PAPI) and tris-(isocyanatophenyl) thiophosphate (Desmodur).

Additional isocyanate components can be prepared by reacting an excess of a diisocyanate as described above with a suitable hydroxyl component, such as a polyol as described above or an R_f -glycol as described herein, or combination thereof, to obtain an isocyanate-terminated prepolymer.

In addition to the polyisocyanate, useful urethane compositions can be obtained from the aliphatic and aromatic monoisocyanates. The low molecular weight urethane compositions obtained by reacting an R_f -glycol with a monoisocyanate are useful to impart soil and mold-release properties to a variety of natural and synthetic polymers.

Some useful aromatic monoisocyanates include -2-fluorophenyl isocyanate, 3-fluorophenyl isocyanate, 4-fluorophenyl isocyanate, m-fluorosulfonylphenyl isocyanate, trans-2-phenylcyclopropyl isocyanate, m-tolyl isocyanate, p-tolyl isocyanate, 1,1,1-trifluoro-o-tolyl isocyanate, 1,1,1-trifluoro-m-tolyl isocyanate, p-bromophenyl isocyanate, 2,5-dimethylphenyl isocyanate, o-ethoxyphenyl isocyanate, p-ethoxyphenyl isocyanate, o-methoxyphenyl isocyanate, m-methoxyphenyl isocyanate, p-methoxyphenyl isocyanate, 1-naphthyl isocyanate, o-nitrophenyl isocyanate, m-nitrophenyl isocyanate, p-nitrophenyl isocyanate, p-phenylazophenyl isocyanate, o-tolyl isocyanate.

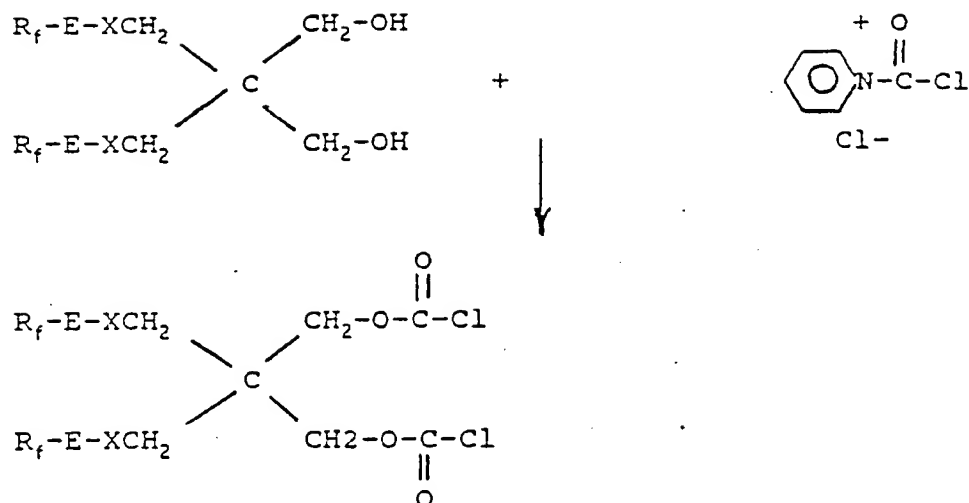
Useful aliphatic or cycloaliphatic monoisocyanates include such alkyl isocyanates of 1 to 16 carbon atoms as methyl isocyanate, ethyl isocyanate, n-propyl isocyanate, n-butyl isocyanate, t-butyl isocyanate, hexyl isocyanate, octyl isocyanate, 2-dodecyl isocyanate, octadecyl isocyanate, hexadecyl isocyanate and mixtures thereof, as well as cyclohexyl isocyanate.

m-Isopropenyldimethyl benzyl isocyanate (TMI) isocyanate-terminated prepolymers typically having a molecular weight of from 200 to about 4000 can be prepared by reacting an excess of an isocyanate component with a polyhydroxy component. The isocyanate component can be a diisocyanate or polyisocyanate as previously described, or can be a low molecular weight isocyanate-terminated prepolymer.

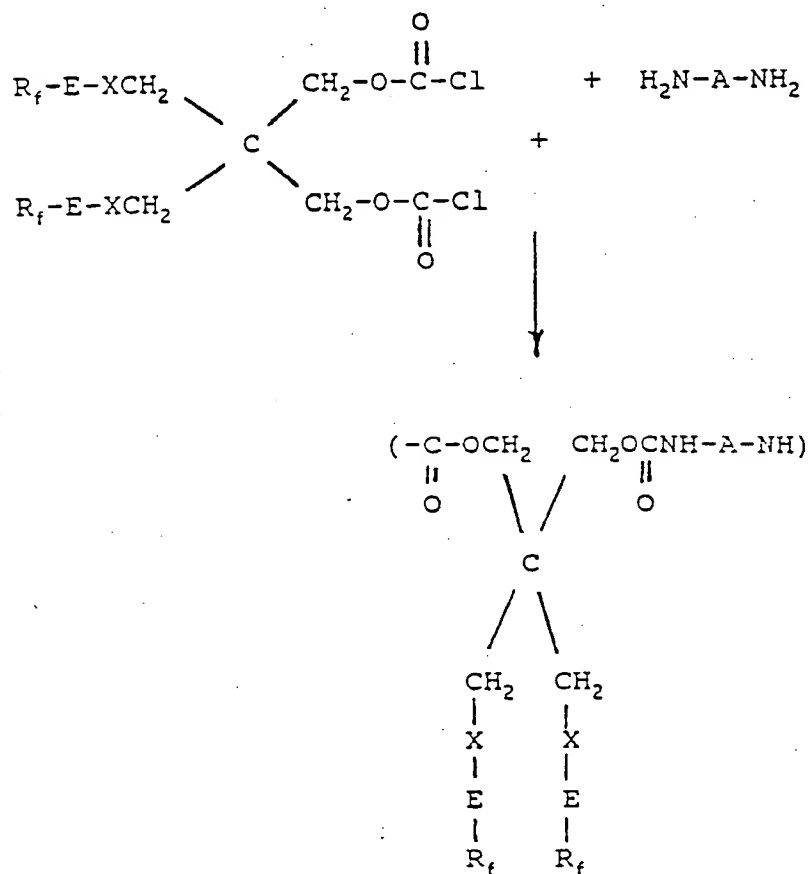
The hydroxy component can be one or more of a polyol, polyester, polyether, polycarbonate and R_f -glycol, all as described previously.

It can be seen that the properties of ultimate urethane compositions can be modified by appropriate modifications in the compositions of the prepolymers.

In addition to the formation of the urethane compositions described above, the R_f -glycols described herein can be converted to the corresponding bischloroformate by treatment with chlorocarbonyl pyridinium chloride:



which in turn can be reacted with an appropriate amine to yield a urethane composition:



where

A is a divalent organic radical as previously described.

The reaction between the isocyanate component and the hydroxyl component can be carried out in bulk, i.e., without solvent, or in the presence of non-reactive, anhydrous, organic solvents. Solvent media in which the reaction can be carried out include ketones, such as acetone, methyl ether ketone and methyl isobutyl ketone; esters such as ethyl acetate, isopropyl acetate, butyl acetate, 2-ethylhexyl acetate; hydrocarbons such as hexane, heptane, octane and higher homologs, cyclohexane, benzene, toluene, xylene or blends of aliphatic, cycloaliphatic and aromatic hydrocarbons or aprotic solvents such as N-methylpyrrolidine; it is also possible to employ ethers, both aliphatic and alicyclic including di-n-propyl ether, di-butyl ether, tetrahydrofuran and the diethers of polyalkylene oxides. In addition, chlorinated solvents such as

1,1,1-tri-chloroethane, dichloroethyl ether, ethylene dichloride, perchloroethylene and carbon tetrachloride can be used.

Among the solvents listed, the water miscible solvents such as acetone and methyl ethyl ketone are most important since they allow conversions of R_1 -urethanes into water soluble R_1 -urethanes as previously described.

In all cases, the solvents should be anhydrous to avoid urea formation.

The reaction can, if desired, be catalyzed and those catalysts conventionally employed in the urethane art are useful herein. Useful catalysts fall principally in two groups -

a. amino compounds and other bases: triethylamine and other trialkylamines, triethylenediamine, 1,4-diaza-2,2,2-bicyclooctane, N-(lower)alkylmorpholines, N,N,N',N'-tetra-methylethylenediamine, N,N,N',N'-tetramethyl-1,3-butanediamine, substituted piperazines, dialkylalkanolamines, benzyltrimethylammonium chloride and

b. organometallic and inorganic compounds: cobalt naphthenate, stannous chloride, stannous octoate, stannous oleate, dimethyl tin dichloride, di-n-butyltin dilaurylmercaptide, tetra-n-butyl-tin, trimethyl-tin hydroxide, di-n-butyltin dilaurate.

Such catalysts may be used singly or in combination with each other. Beneficial synergistic catalysis may occur when combinations are used.

While it is possible to carry out the reaction without the use of a catalyst, it is preferable for reasons of economy and to assure a complete reaction, to utilize one or more catalysts as listed in amounts ranging from 0.001 to 1% based on the weight of the reactants. It is similarly advantageous to carry out the urethane synthesis at elevated temperature, usually between room temperature and 120°C and preferably at 60°C to 80°C to obtain a complete reaction between 0.5 to 8 hours reaction time.

The reaction can be easily followed by titration of the isocyanate group or by IR analysis.

The determination of the critical surface tension in dynes per centimeter shows that the free surface energy of a polyurethane is lowered if the novel R_1 -glycols are incorporated into the urethane chain.

The critical surface tensions are determined by contact angle measurements as described by W. Zisman, Contact Angles, Advances in Chemistry, No. 43, ACS Publications, Washington, D.C., 1964.

The usefulness of the polyurethane compositions is, conveniently shown by measuring the oil, water and soil repellency ratings of substrates such as fabrics, paper, leather, etc. which are treated with solutions or emulsions of the novel urethane compositions.

As already indicated, the urethane compositions of the invention are highly effective for imparting oil and water repellent properties to substrates to which they are applied and coatings of these polymers may be prepared by any of the well-known techniques. When prepared by bulk or suspension polymerization techniques, these urethane compositions can be applied, for example, from a dilute solution in suitable a solvent such as the fluoroalkanes, fluorochloroalkanes, fluoroalkanoic acids, chlorinated alkanes or aromatics, hydrocarbon aromatics, ketones, ester and others. Concentrations of the fluorinated polymer in the solvent can be adjusted to provide an amount of urethane composition deposited on the substrate sufficient to provide oil and water repellency. This amounts typically to a deposit of from 0.01 to 10%, preferably from 0.1 to 1%, of urethane composition, based on the weight of substrate. If the urethane composition is obtained as an aqueous latex or emulsion, the system can be diluted with water or other appropriate diluent to similarly provide an amount of urethane ranging from 0.01 to 10% of the weight of substrate deposited thereon.

The urethane solution or latex may be applied by any of the known techniques such as by dipping, spraying, brushing, padding, roll coating or by any desired combination of such techniques. The optimum method of application will depend principally on the type of substrate being coated.

Coatings of the urethane compositions of the invention may be applied to any desired substrate, porous or non-porous. They are particularly suited for application to porous materials such as textiles, leather, paper, wood, masonry, unglazed porcelain and the like to provide valuable oil and water repellency properties. However, they may also be applied to non-porous materials such as metals, plastics, glass, painted surfaces and the like to provide similar oil and water repellency properties. More specifically the urethane compositions of the invention act as levelling, wetting and spreading agents in formulations designed for application to floors, furniture and automobiles. In such applications a protective oil and water repellent film is left on the treated object after the removal of the bulk of the material. Such levelling, wetting, spreading and film forming properties are also useful in

a. formulations for cleaning glass and other hard, non-porous materials

b. hair care products such as rinses, shampoos and hair sprays.

c. paint, stain and varnish formulations for application to wood, masonry and ceramics.

In the treatment of paper the urethane compositions may be present as an ingredient in a wax, starch, casein, elastomer, or wet strength resin formulation. Aqueous emulsions of the urethane compositions are especially useful in the treatment of paper. By mixing the urethane compositions in an aqueous or oil type paint formulation, it may be applied effectively to unpainted asbestos siding, wood, metal and masonry. In the treatment of floors and tile surfaces and like substrates, and urethane compositions may be applied by their incorporation in an emulsion or solution.

Because of the ability of the surfaces treated with these urethane compositions to withstand abrasive action, the advantages incident to the repellency to oil and water and their resistance to soiling imparted by coating them with the urethane compositions of this invention, preferred classes of articles to be treated are

papers and textiles. Illustrative papers are carbonizing tissue, wallpaper, asphalt laminates, liner board, cardboard and papers derived from synthetic fibers.

For application to textile materials such as fabrics woven and non-woven, fibers, films, yarns, cut staple, thread etc. or articles made from fabrics, fibers, films, yarns, etc. the urethane compositions of the invention are preferably prepared as aqueous latices or emulsions which are then diluted, preferably with water and applied to the textiles from pad baths which may contain other treating materials. In accordance with this technique, the fabric or the textile material is passed through the bath, passed through squeeze rolls adjusted to leave the desired amount of the latex on the fabric, dried at a temperature of about 25°C to 125°C and then cured in a curing oven at a temperature in the range of from 20°C to 195°C for 0.2 to 20 minutes. The weight of urethane composition deposited on the fabric may range from 0.01 to 10% of the weight of fabric. Preferably, very small amounts are used in the range of 0.1 to 1%, often from 0.1 to 0.5% to give high degrees of water and oil repellency. Any types of textile materials, such as cotton, wool, fiber glass, silk, regenerated cellulose, cellulose esters, cellulose ethers, polyesters, polyamides, polyolefins, polyacrylonitrile, polyacrylic esters, inorganic fibers, etc. either along or blended in any combination may be successfully coated with the urethane compositions of the invention. The resulting textile material will be found to be repellent to water and oil, and the textile material will retain its resistance to such agents even after many launderings and dry cleanings.

It will be often advantageous to use the urethane compositions of the invention in combination with conventional finishes, such as mildew preventatives, moth resisting agents, crease resistant resins, lubricants, softeners, fat liquors, sizes, flame retardants, antistatic agents, dye fixatives and water repellents.

TEST METHODS

The AATCC water spray test rating was determined according to Standard Test method 22-1985 of the American Association of Textile Chemists and Colorists, Volume 61, 1986 (also designated ASTM-D-583-58). Ratings are given from 0 (minimum) to 100 (maximum).

The AATCC Oil Rating was determined according to Standard Test method 118-1983 of the American Association of Textile Chemists and Colorists. Ratings are given from 0 (minimum) to 8 (maximum). A commonly accepted level of repellency for oil repellent fabrics in the United States is an oil repellency of 4.

All mentioned AATCC Tests are listed in the Technical manual of the American Association of Textile Chemists and Colorists, volume 61, edition 1986.

Polymers prepared in water or a water-solvent mixture or a solvent which is water-miscible may be applied to polyester-cotton twill by padding from an aqueous pad bath containing also the following permanent press resins, catalyst and additives (so-called permanent press recipe):

After the padding, the fabric is dried at 100°C for 2 minutes and cured at 163°C for 5 minutes.

The invention described above is illustrated by the following examples.

EXAMPLE 1



1,1,2,2-Tetrahydroperfluorooctanethiol (176.7 g, 0.465 mol) and dibromoneopentyl glycol (60.8 g, 0.232 mol) are reacted under nitrogen with potassium carbonate (64.3 g, 0.465 mol) and 2-pentanone (53.2 g) as the solvent. The reaction is carried out at 105°C for 16 hours. The product is stirred at 70°C and washed once with 350 ml distilled water. Residual water is removed as an azeotrope and the remaining solvent is evaporated under house vacuum. The diol (187.5 g, 94% of theory) is of 89% purity by GLC. The crude product is recrystallized twice from 500 g toluene to give a final product 95% pure by GLC, m.p. 71.5-73.5°C. Analysis for $\text{C}_{21}\text{H}_{18}\text{F}_{26}\text{SO}_2\text{S}_2$:

Calculated: C, 29.3; H, 2.1; F, 57.4; S, 7.5.

Found: C, 29.4; H, 2.0; F, 56.6; S, 7.7.

EXAMPLE 2

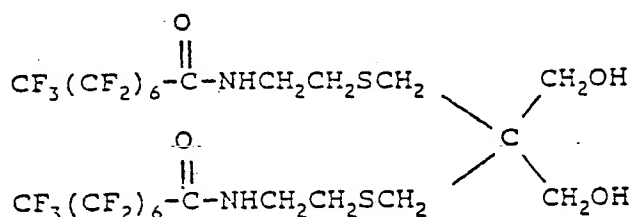


1,1,2,2-Tetrahydroperfluorooctanethiol (111.6 g, 0.232 mol) and dibromoneopentyl glycol (30.4 g, 0.116

mol) are reacted under nitrogen with potassium carbonate (32.1 g, 0.232 mol) and 2-pentanone (30.7 g) as the solvent. The reaction is carried out at 100°C for 17 hours and the by-product salts (KBr) is removed by a hot gravity filtration. The crude product is crystallized and vacuum filtered giving 101.1 g (82% of theory) of a pale yellow solid, 77% pure by GLC. The diol is recrystallized twice from toluene (400 g) and once from carbon tetrachloride (400 g) to give a final product of 96% purity, m.p. 90-92.5°C. Analysis for C₂₅H₁₈F₃₄O₂S₂:

Calculated: C, 28.3; H, 1.7; F, 60.9; S, 6.1.
Found: C, 28.3; H, 1.9; F, 60.9; S, 6.5.

EXAMPLE 3

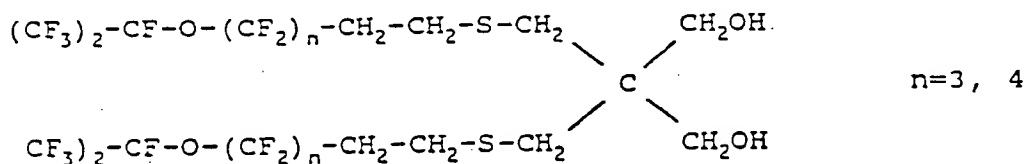


N-[2-Mercaptoethyl]-perfluoro-octanamide (45.3 g, 0.10 mol) and dibromoneopentyl glycol (12.5 g, 0.048 mol) are reacted under nitrogen with potassium carbonate (13.2 g, 0.10 mol) and 2-pentanone (14.0 g) as the solvent. The reaction is run at 103°C for 19.5 hours. The reaction mixture is stirred at 75°C and washed twice with 80 g distilled water. The solvent is evaporated below 115°C to yield a brown product which is transferred as a brittle solid. This solid is recrystallized from toluene and then from ethanol to yield a white solid (32 g, 64% of theory) which is 93% pure by GLC. NMR showed proton resonances at 2.72 ppm, 4 protons, S-CH₂C; 2.79 ppm, 4 protons, CH₂SCH₂; 3.60 ppm, 4 protons, NH-CH₂; 3.65 ppm, 4 protons, CH₂-OH; 6.97 ppm, 1 proton, N-H.

Analysis for C₂₅H₂₀F₃₀N₂O₄S₂:

Calculated: C, 28.7; H, 1.9; N, 2.7; F, 54.5; S, 6.1.
Found: C, 28.5; H, 1.7; N, 2.8; F, 54.6; S, 6.3.

EXAMPLE 4



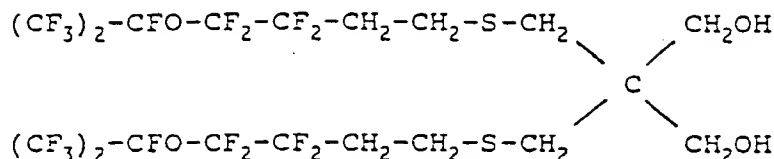
1-Heptafluoroisopropoxy-1,1,2,2-tetrahydroperfluoroalkanethiol (consisting of 73% n=3 homolog and 27% n=4 homolog) (46.0 m, 0.080 mol) and dibromoneopentyl glycol (10.5 m, 0.040 mol) are reacted under nitrogen with potassium carbonate (11.1 m, 0.080 mol) and 2-pentanone (13.0 g) as the solvent. The reaction is carried out at 103°C for 18 hours. The reaction mixture is stirred at 75°C and washed twice with 80 g distilled water. The solvent is evaporated below 115°C to yield a yellow oil which solidifies upon cooling to a waxy yellow gel (32.3 g, 64% of theory). The product is 87% pure by GLC. NMR confirmed structure with the general proton assignments are the same as in example 1.

Analysis for product (n=3, 4)

Calculated: C, 28.1; H, 2.0; F, 55.9; S, 7.0.

Found: C, 27.7; H, 1.6; F, 59.5; S, 5.3.

EXAMPLE 5



4-Heptafluoroisopropoxy-1,1,2,2-tetrahydroperfluorobutanethiol (43.7 g , 0.13 mol) and dibromoneopentyl glycol (16.5 g , 0.06 mol) are reacted under nitrogen with potassium carbonate (17.4 g , 0.13 mol) and 2-pentanone (15.0 g) as the solvent. The reaction is carried out at 103°C for 19 hours. The product is stirred at 75°C and washed twice with 80 ml. distilled water. The solvent is evaporated below 115°C and the product transferred hot as a yellow oil which remains fluid. The diol (31.4 g , 63% of theory) is of 87% purity by GLC. NMR showed proton resonances at 2.01 ppm, 2 protons, -OH; 2.1-3.0 ppm, 8 protons, CF₂CH₂CH₂S; 2.70 ppm, 4 protons, SCH₂C; 3.66 ppm, 4 protons -(CH₂-OH)₂.

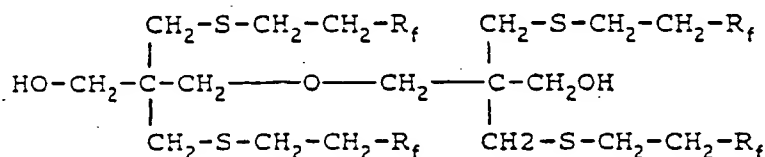
Analysis for product $C_{19}H_{18}F_{22}O_4S_2$:

Calculated: C, 28.8; H, 2.3; F, 52.7; S, 8.1.

Found: C, 28.9; H, 2.4; F, 50.4; S, 7.6.

EXAMPLE 6

Tetrakis-(1,1,2,2-tetrahydroperfluoroalkanethio)dipentaerythritol



1,1,2,2-Tetrahydroperfluoroalkaneithiol* (139.6 g, 0.28 mol) and sym-tetrabromodipentaerythritol (35.0 g, 0.07 mol) are reacted under nitrogen with potassium carbonate (38.7 g, 0.28 mol) and 2-pentanone (60.0 g) as the solvent. The reaction is run at 103°C for 19.5 hours, and is then allowed to cool and washed twice with 160 g distilled water at 75°C. The water layer is removed and the remaining water/2-pentanone mixture azeotroped below 115°C. The product is isolated as a brittle, yellowish-brown solid and crystallized from toluene to yield a pale yellow solid (127.4 g, 86% of theory) m.p. 79.7-87.4°C NMR shows proton resonances at 1.93 ppm, 2 protons, O-H; 2.40 ppm, 8 protons, $R_1\text{-CH}_2$; 2.63-2.90 ppm, 16 protons, $\text{CH}_2\text{-S-CH}_2$; 3.43 ppm, 4 protons, $\text{CH}_2\text{-O-CH}_2$; 3.65 ppm, 4 protons, $\text{CH}_2\text{-OH}$.

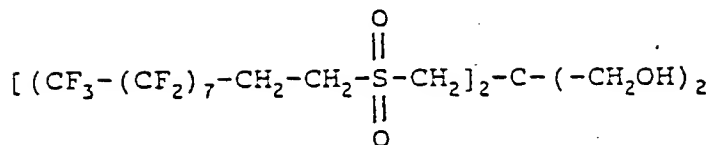
Analysis for % OH:

Calculated = 1.58%

Found = 1.56%

- R_f distribution C₈ / C₁₀ - 88.90%/6.40%

EXAMPLE 7

2,2-Bis(1,1,2,2-tetrahydroperfluorodecylsulfonylmethylene)-1,3-propanediol

2,2-Bis-(1,1,2,2-tetrahydroperfluorodecylthiomethylene)-1,3-propanediol (25.0 g, 0.024 mol) is dissolved in glacial acetic acid (38.09 g, 0.63 mol) and warmed to 40°C. Hydrogen peroxide (5.4 g, 30%) is added and the mixture is stirred for hour. The reaction mixture is then heated to 100°C and additional hydrogen peroxide (10.6 g, 30%) is added. The product precipitates as a yellowish-white solid (24.6 g, 95% of theory) and is crystallized from isopropyl acetate to yield a white solid, m.p. 165-168°C which was 92% pure by GLC. NMR shows proton resonance at 2.82 ppm, 4 protons, CF₂-CH₂; 3.63 ppm, 4 protons, CH₂-SO₂-CH₂; 3.93 ppm, 4 protons, SO₂-CH₂-C; 4.25 ppm, 4 protons, -C-CH₂OH.

Analysis for C₂₅H₁₈F₃₄O₆S₂;

Calculated: C, 26.70; H, 1.6; F, 57.44; S, 5.70.

Found: C, 26.60; H, 1.5; F, 54.50; S, 6.30.

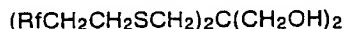
EXAMPLE 8

Reaction Products of (RfCH₂CH₂SCH₂)₂-C-(CH₂OH)₂ and γ-Caprolactone

2,2-Bis(1,1,2,2-tetrahydroperfluoroalkanethiomethylene)-1,3-propanediol* (108.5 g, 0.10 mol) and S-caprolactone (45.7 g, 0.40 mol) are reacted under nitrogen at 75°C for 1.5 hours in the presence of a catalytic amount of stannous octanoate. The reaction is then completed at 140°C for 3 hours. The product is a soft waxy yellow solid of mp 28-34°C. Comparative gel permeation chromatography (styrogel column, THF mobile phase) of the starting material and the product shows an increase in the weight average molecular weight from 1091 to 2508.

* R_f distribution C₆F₁₃/C₆F₁₇-90%/88%

EXAMPLE 9



1,1,2,2-Tetrahydroperfluoroalkanethiol* (215.8 g, 0.47 mmol) and dibromoneopentyl glycol (60.78 g, 0.23 mmol) are reacted under nitrogen with potassium carbonate (64.3 g, 0.465 mmol) and 2-pentanone (54 g) as the solvent. The reaction is carried out at 105°C for 17 hours. The product is stirred at 70°C and washed once with 350 ml distilled water. Residual water is removed as an azeotrope and the remaining solvent is evaporated under house vacuum. The diol (228.3 g, 92% of theory) is of 89% purity by GLC.

* R_f-distribution % C₄/C₆/C₈/C₁₀ - 0.1%/9/88/1.2

EXAMPLE 10

Dibromoneopentyl glycol is reacted with 1,1,2,2-tetrahydroperfluoro-decyl amine in triethylamine as solvent at 80-100°C. The reaction is carried out for 12 hours and the product is worked up by acidification and precipitation into water. The powder is collected by filtration and dried to yield 2,2-Bis-(1,1,2,2-tetrahydroperfluorodecyliminomethylene)-1,3-propanediol.

EXAMPLE 11

1,1,2,2-Tetrahydroperfluorooctanol (10.9 g , 0.03 mol), 3,3-(bromomethyl)oxetane (3.9 g , 0.016 mol), potassium hydroxide (1.7g , 0.03 mol), and 15 g of diethylene glycol dimethyl ether are reacted in the presence of approximately 1.0 g of 18 Crown-6 in a 50 ml flask with gentle stirring for 72 hours at room temperature. Analysis by gas chromatography indicates that the solution contains 36% monoadduct, 64% diadduct. 2.5 g of solids are filtered off and GC/MS identification confirms the presence of the desired 3,3-(1,1,2,2-tetrahydroperfluorooctyloxy)oxetane and 3-(1,1,2,2-tetrahydroperfluorooctyloxy)-3-bromooxetane.

Identification is made by GC/MS (E.I. Mode) after derivatization with N,O-bis-(trimethylsilyl)trifluoroacetamide. Two major components are observed:

(Diadduct)

MS: m/z 810 absent, 780 (M-CH₂O), 417 b.p., (M-C₆F₁₃CH₂CH₂O), 403 (M-C₆F₁₃CH₂CH₂OCH₂).

MS (CI) 811 (M + 1)⁺

(Monoadduct)

MS: m/z 526 absent, 447 (M-Br), 417 b.p. (M-CH₂OBr).

MS (CI) 527 (M + 1)⁺

EXAMPLE 12

In a similar fashion to the previous example, 1,1,2,2-tetrahydro-perfluorodecanol is reacted at 100°C for 2-3 hours to give a mixture of monoadduct=40% and diadduct=60%.

EXAMPLE 13

The hydrolysis of the mixture of mono- and di- C₆F₁₃CH₂CH₂O-oxetanes obtained in Example 11 is accomplished by reacting the solution of the subject oxetanes in diethylene glycol dimethyl ether with 10 gm water containing 1 g concentrated sulfuric acid, at reflux for 48 hours. The product separates into two layers and the bottom layer is separated off and washed once with dilute potassium carbonate and twice with water. Evaporation of 10 grams of product at 220°C/1mm Hg with a Kugelrohr affords 8 grams of an oil. By gas chromatography the product contains 79% mono- and 21% diadduct diols, as well as a considerable amount of byproducts derived from the solvent.

(Diadduct)

MS (C.I.) = 29 (M + H)⁺

MS (E.I.) = m/z 828 (absent), 417 b.p. (C₁₂H₁₀OF₁₃), 403 (C₁₁H₈O₁₃).

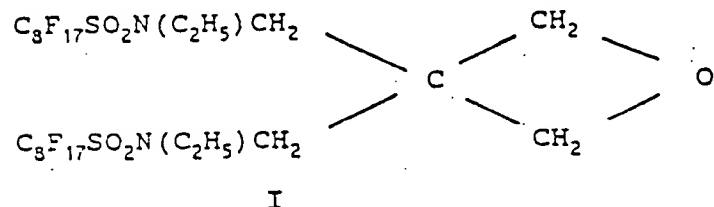
(Monoadduct)

MS (C.I.) = 545 (M + H)⁺

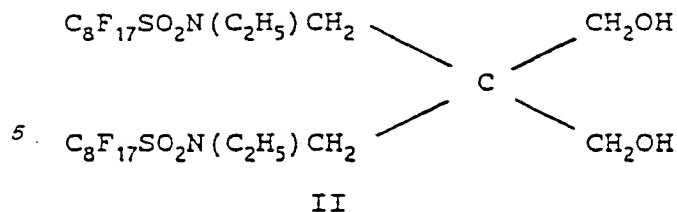
MS (E.I.) = m/z 544 (absent), 447 (-H₂O, Br), 429 (-2H₂O, Br), 417 (-CH₂O, Br, H₂O), 71 b.p. (C₄H₇O).

Example 14

A mixture of C₈F₁₇SO₂N(C₂H₅)H (20.0g, 0.047mol), 87% potassium hydroxide (3.0g, 0.047 mol), diglyme (60.0g) and 18-Crown-6 ether (0.1g) is heated under nitrogen at 100 - 105°C for 30 minutes. The 3,3-bis-(bromomethyl)-oxetane (5.7g, 0.0236 mol) is then added and the entire reaction mass is heated at 100°C for 13 hours. The salts are removed upon filtration. Diglyme is removed at 80°C under vacuum. Proton NMR shows the presence of I in the product.



The oxetane of I can be opened to yield the diol II,



10 according to the procedure described in example 13.

Examples 15 to 24

15 Using the methods described and by techniques similar to Examples 1-14, the following additional sulfido-
and sulfone diols are prepared.

Ex.	Thiol	Perfluoroalkyl Terminated Neopentyl Glycol
20	15 $\text{CF}_3\text{CF}_2\text{CH}_2\text{SH}$	$(\text{CF}_3\text{CF}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})_2$
	16 $\text{C}_6\text{F}_{13}(\text{CH}_2)_4\text{SH}$	$(\text{C}_6\text{F}_{13}(\text{CH}_2)_4\text{SO}_2\text{CH}_2)_2\text{C}(\text{CH}_2\text{OH})_2$
	17 $\text{C}_8\text{F}_{17}\text{CH}_2\text{CH}_2\text{CH}_2\text{SH}$	$(\text{C}_8\text{F}_{17}\text{CH}_2\text{CH}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})_2$
	18 $\text{C}_8\text{F}_{17}\text{CH}_2\text{CH}_2\text{N}(\text{CH}_3)\text{CH}_2\text{CH}_2\text{CH}_2\text{SH}$	$(\text{C}_8\text{F}_{17}\text{CH}_2\text{CH}_2\text{N}(\text{CH}_3)\text{CH}_2\text{CH}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})_2$
	19 $\text{C}_8\text{F}_{17}\text{SO}_2\text{NHCH}_2\text{CH}_2\text{SH}$	$(\text{C}_8\text{F}_{17}\text{SO}_2\text{NHCH}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})_2$
25	20 $\text{C}_8\text{F}_{17}\text{CH}_2\text{CH}_2\text{SO}_2\text{NHCH}_2\text{CH}_2\text{SH}$	$(\text{C}_8\text{F}_{17}\text{CH}_2\text{CH}_2\text{SO}_2\text{NHCH}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})_2$
	21 $\text{C}_7\text{F}_{15}\text{CONHCH}_2\text{CH}_2\text{SH}$	$(\text{C}_7\text{F}_{15}\text{CONHCH}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})_2$
	22 $\text{CF}_3\text{CF}_2\text{CH}_2\text{CH}_2\text{SH}$	$\text{HO}[\text{CH}_2\text{C}(\text{CH}_2\text{SCH}_2\text{CH}_2\text{CF}_2\text{CF}_3)_2\text{CH}_2\text{O}]_2\text{-H}$
	23 $\text{C}_8\text{F}_{17}\text{CH}_2\text{CH}_2\text{SH}$	$\text{HO}[\text{CH}_2\text{C}(\text{CH}_2\text{SO}_2\text{CH}_2\text{CH}_2\text{C}_8\text{F}_{17})_2\text{CH}_2\text{O}]_2\text{-H}$
	30	24 Reaction product of $(\text{R}_1\text{CH}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})_2$ and Gamma-caprolactone

EXAMPLE 25

Stability Comparisons of Bis-perfluoroalkylthio-Diols

40 Thermogravimetric analyses are run on the diols of Examples 2, 6, 7 and 8.
The instrument is run at 10°C/minute to either 350°C or 500 C with 100 ml/min N₂ or in air.

TGA (Wt. Loss)

Temperature (°C) of Indicated Weight Loss

Diol of Example	DSC (Tm)	Under Nitrogen			Under Air		
		Init.	10%	50%	Init.	10%	50%
2	79-86	150	251	295	150	237	285
7	165-168	195	286	328	185	260	309
8	28-34	150	293	340	150	265	316
6	79-87	180	315	356	185	285	335

These results indicate the thermal stability of the subject bis-perfluoroalkylthio-neopentyl type diols. The subject diols as exemplified above all are appreciably stable to greater than 280°C in nitrogen, even to 260°C in air. The weight loss of the diol of Example 2 is appreciable by TGA because of volatilization, not decomposition.

EXAMPLE 26

The diol of Example 9 (78.2 g, 0.073 mmol) is predried azeotropically with 1,1,1-trichloroethane. It is then
65 reacted with 3,3,4-trimethylhexane-1,6-diisocyanate (11.54 g, .055 mol) under nitrogen in the presence of a

catalytic amount of stannous octanoate (0.22 g) with 1,1,1-trichloroethane (135 g) as the solvent. The reaction mixture is heated at reflux for one hour at which time the $\text{N}=\text{C}=\text{O}$ infrared band at 2270cm^{-1} is absent. Then, dimer acid diisocyanate (27.7 g, 46 mmol) is added along with N-methyldiethanolamine (3.4 g, 28 mmol) and 1,1,1-trichloroethane (46.6 g).

The mixture is kept at reflux for two hours and the reaction is again judged to be complete by the disappearance of the $\text{N}=\text{C}=\text{O}$ infrared band. Molecular weight can be determined by gel permeation chromatography.

EXAMPLE 27

The diol of Example 6 (30.0 g, 0.014 mol) is predried azeotropically with isopropyl acetate and is then reacted with 3,3,4-trimethylhexane-1,6-diisocyanate (2.19 g, 0.01 mol) under nitrogen in the presence of a catalytic amount of stannous octanoate (0.084 g) using isopropyl acetate (49.0 g) as the solvent. The reaction mixture is heated to 80°C for 30 minutes and completeness of reaction is indicated by the absence of the $\text{N}=\text{C}=\text{O}$ infrared band at 2270 cm^{-1} . Then, dimer acid diisocyanate (10.36 g, 0.017 mol) is added along with N-methyldiethanolamine (1.67 g, 0.014 mol) and isopropyl acetate (18.0 g). The mixture is heated to 80°C for one hour until the reaction is judged to be complete by the absence of the $\text{N}=\text{C}=\text{O}$ infrared band. Molecular weight can be determined by gel permeation chromatography.

EXAMPLE 28

The diol of Example 6 (30.0 g, 0.014 mmol) is predried azeotropically with isopropyl acetate and is then reacted with 3,3,4-trimethylhexane-1,6-diisocyanate (1.45 g., 0.007 mmol) under nitrogen in the presence of a catalytic amount of stannous octanoate (0.084 g), using isopropyl acetate (47.0 g) as the solvent. The reaction mixture is heated at 80°C for 30 minutes and reaction is complete as indicated by the absence of the $\text{N}=\text{C}=\text{O}$ infrared band at 2270 cm^{-1} . Then dimer acid diisocyanate (10.4 g, 17 mmole) is added along with N-methyldiethanolamine (1.25 g, 10 mmole) and isopropyl acetate (17.40 g). The mixture is heated at 80°C for 1 hour and is again judged to be complete by the absence of the $\text{N}=\text{C}=\text{O}$ infrared band.

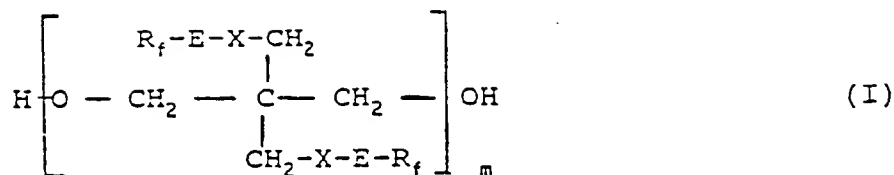
Example 29

The polyurethane products of Examples 26 and 28 are applied by pad application with 1,1,1-trichloroethane on both 65/35 polyester/cotton poplin and 100 % nylon taffeta in such a way that 0.06 % F is applied to the polyester/cotton and 0.04 % F to the nylon. The products are evaluated after air drying and curing in a hot air oven at $> 150^{\circ}\text{C}$ for 3 minutes. The products prepared from the subject diols have significantly improved water repellency over the prior art diol.

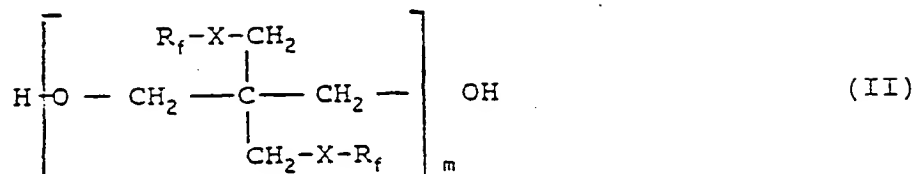
Product of Example	65/35 polyester/cotton at 0.06% F		100 nylon taffeta at 0.04% F
	Air Dried/Oven Cured AATCC Oil	AATCC Spray	Air Dried AATCC Spray
21	-	-	100
23	4	100	100

Claims

1. A compound of formulas I or II



OR



wherein R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of -NR-, -O-, -S-, -SO₂-, -COO-, -OOC-, -CONR-, -NRCO-, -SO₂NR-, and -NRSO₂-, or said alkylene terminated at the R_f end with -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and for formula I, X is -S-, -O-, -SO₂-, or -NR-, and for formula II, X is -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and where R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms, and m is 1, 2 or 3.

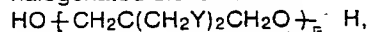
2. A compound of formula I or II according to Claim 1 wherein R_f is a straight or branched chain perfluoroalkyl of 2 to 12 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, and m is 1 to 2.

3. A compound of formula I according to Claim 1 where R_f is perfluoroalkyl of 2 to 12 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, CH₂CH₂OCH₂CH₂-, or -SO₂NHCH₂CH₂-, X is -S-, -SO₂- or -O-, and m is 1 or 2.

4. A compound of formula I according to Claim 1 wherein R_f is perfluoroalkyl of 6 to 12 carbon atoms, E is ethylene, X is -S-, and m is 1.

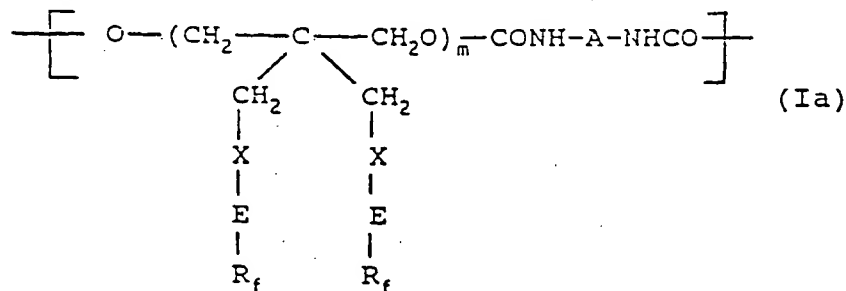
5. A compound of formula I according to Claim 1 wherein R_f is perfluoroalkyl of 6 to 12 carbon atoms, E is ethylene, X is -S-, and m is 2.

6. A process for the preparation of compounds of the formulas I or II, which comprises reacting compounds of formulas $R_f-E-X-H$ or R_f-X-H , wherein R_f , E and X have the meanings given in claim 1, with a halogenated diol of formula

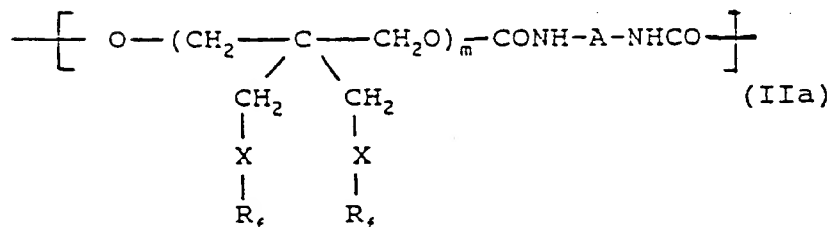


wherein m is 1, 2 or 3 and Y is Cl, Br or I.

7. A linear polyurethane having recurring structural units of the formula Ia or IIa:



OR

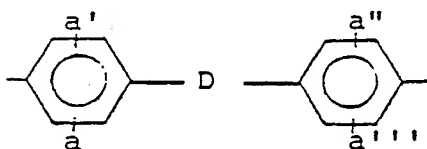


wherein R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl

substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of -NR-, -O-, -S-, -SO₂-, -COO-, -OOC-, -CONR-, -NRCO-, -SO₂NR-, and -NRSO₂-, or terminated at the R_f end with -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and for formula Ia, X is -S-, -O-, -SO₂-, or -NR-, and for formula IIa, X is -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and where R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms, and m is 1, 2 or 3, and A is a divalent aliphatic, cycloaliphatic or aromatic group.

8. A polyurethane according to claim 7 wherein A is alkylene of 2 to 16 carbon atoms, cycloalkylene of 6 to 15 carbon atoms, phenylene, naphthylene or said phenylene or said naphthylene substituted by one or two alkyl of 1 to 4 carbon atoms, by alkoxy of 1 to 4 carbon atoms, by chloro, by bromo or by nitro, or A is a group of the formula



where D is a direct bond, oxygen, methylene or ethylene, and a, a', a'' and a''' each independently are hydrogen, alkyl of 1 to 4 carbon atoms, alkoxy of 1 to 4 carbon atoms, chloro or bromo.

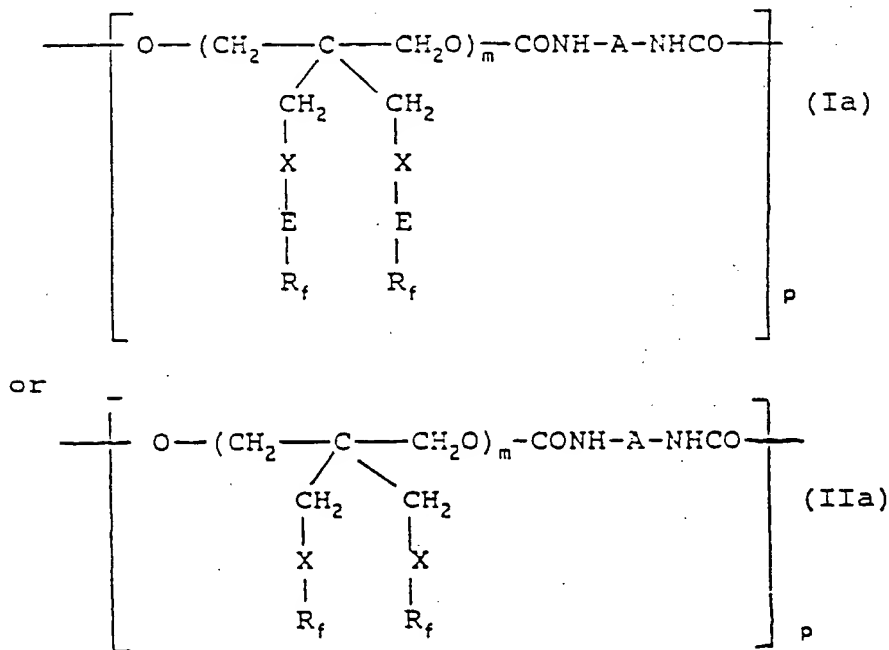
9. A polyurethane according to claim 7 wherein A is 3,3,4-trimethyl-1,16-hexanediyl or 1,3,3-trimethyl- α , α ,5-hexahydrobenzylene.

10. A polyurethane according to claim 7 where R_f is perfluoroalkyl of 2 to 12 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, -CH₂CH₂OCH₂CH₂-, or -SO₂NHCH₂CH₂-, X is -S-, -SO₂- or -O-, and m is 1 or 2.

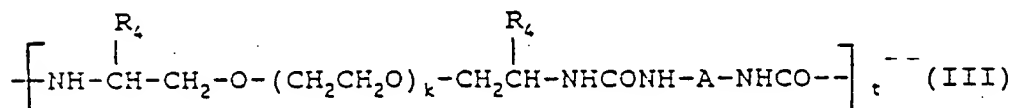
11. A linear block copolymer having soil-release properties which contains, based on the total block copolymer,

(a) 15-70% by weight of recurring structural units of blocks Ia, IIa or a mixture thereof,

(b) 85 to 30% by weight of recurring structural units of block III, wherein the blocks are connected to one another via urea linkages, blocks Ia or IIa having recurring structural units of the formula



block III having recurring structural units of the formula:



wherein R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of -NR-, -O-, -S-, -SO₂-, -COO-, -OOC-, -CONR-, -NRCO-, -SO₂NR-, and -NRSO₂-, or terminated at the R_f end with -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and for formula Ia, X is -S-, -O-, -SO₂-, or -NR-, and for formula IIa, X is -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and where R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms, and m is 1, 2 or 3.

A is a divalent aliphatic, cycloaliphatic or aromatic group,

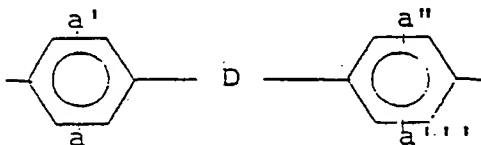
R is hydrogen or methyl,

k is 8 to 100, and

p and t are integers and are independently 1 to 5.

12. A block copolymer according to claim 11 wherein R_f is perfluoroalkyl of 2 to 12 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, -CH₂CH₂OCH₂CH₂-, or -SO₂NHCH₂CH₂-, X is -S-, -SO₂- or -O-, and m is 1 or 2.

13. A block copolymer according to claim 11 wherein A is alkylene of 2 to 16 carbon atoms, cycloalkylene of 6 to 15 carbon atoms, phenylene, naphthylene or said naphthylene substituted by one or two alkyl of 1 to 4 carbon atoms, by alkoxy of 1 to 4 carbon atoms, by chloro, by bromo or by nitro, or A is a group of the formula



where D is a direct bond, oxygen, methylene or ethylene, and a, a', a'' and a''' each independently is hydrogen, alkyl of 1 to 4 carbon atoms, alkoxy of 1 to 4 carbon atoms, chloro or bromo.

14. A block copolymer according to claim 11 wherein A is 3,3,4-trimethyl-1,6-hexanediyl or 1,3,3-trimethyl- α ,5-hexahydrobenzylene.

15. A linear polyester having recurring structural units of the formula Ib or IIb

--[OCH₂C(CH₂X-E- R_f)₂CH₂O]_m-CO-L-CO]- (Ib)

or

--[OCH₂C(CH₂X- R_f)₂CH₂O]_m-CO-L-CO]- (IIb)

wherein R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of -NR-, -O-, -S-, -SO₂-, -COO-, -OOC-, -CONR-, -NRCO-, -SO₂NR-, and -NRSO₂-, or terminated at the R_f end with -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and for formula Ib, X is -S-, -O-, -SO₂-, or -NR-, and for formula IIb,

X is -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and where R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms, and m is 1, 2 or 3.

L is a direct bond, a straight or branched chain alkylene of 1 to 34 carbon atoms, cycloalkylene of 6 to 12 carbon atoms, or arylene of 6 to 16 carbon atoms.

16. A polyester according to claim 15 where R_f is perfluoroalkyl of 2 to 12 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, -CH₂CH₂OCH₂CH₂-, or -SO₂NHCH₂CH₂-, X is -S-, -SO₂- or -O-, and m is 1 or 2.

17. A polyester according to claim 15 wherein L is tetramethylene or o-phenylene.



European Patent
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EUROPEAN SEARCH REPORT

Application Number

EP 89 81 0449

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 4)
A, D	US-A-3 478 116 (K.C. SMELTZ) ---		C 07 C 149/18
A	DE-A-2 702 607 (CIBA-GEIGY) ---		C 07 C 147/02
A	EP-A-0 010 523 (CIBA-GEIGY) -----		C 07 C 147/14
			C 07 C 103/167
			C 07 C 103/34
			TECHNICAL FIELDS SEARCHED (Int. Cl. 4)
			C 07 C 149/00
			C 07 C 147/00
			C 07 C 91/00
			C 07 C 143/00
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 03-10-1989	Examiner VAN GEYT J.J.A.
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